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(54) Title: METHODS AND COMPOSITIONS USEFUL IN PREVENTING EQUINE INFLUENZA (57) Abstract <p>Recombinant vaccines for immunizing horses against equine influenza virus (EIV). The DNA sequences encoding the hemagglutinin (HA) and neuraminidase (NA) glycoproteins from the two strains of EIV currently infective in horses are used to construct vaccinia carried vaccines, to design synthetic peptides for primer and booster administration, and to permit recombinant synthesis of HA and/or NA protein based vaccines. These DNA sequences also provide probes useful for preparing similar vaccines from fresh isolates of new strains generated by genetic drift.</p> <p style="text-align: center;">127</p>		

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METHODS AND COMPOSITIONS USEFUL IN
PREVENTING EQUINE INFLUENZA

Technical Field

5 The invention relates to immunizing horses
against infection by influenza virus. More
particularly, the invention relates to use of
vaccinia-carried immunogens and synthetic peptide
vaccines useful for this purpose.

10

Background Art

Equine influenza is a highly contagious
respiratory infection engendered in horses by equine
influenza virus (EIV). While the disease has low
15 mortality, the economic impact is often great due to
deterioration of the subject's performance (Mumford,
J.A., et al, Equine Vet J (1980) 12:3-9). Current
immunization techniques use inactivated or killed EIV.
These procedures are characterized by undesirable side
20 effects, and the immunity conferred appears in some
cases to last for no more than three or four months. In
short, there is no entirely satisfactory vaccine
available to prevent the spread of this disease.

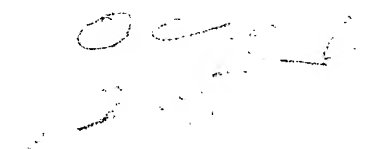
Two different serotypes, which are members of
25 the influenza A myxovirus group, have been identified as
causative agents of this disease: A/equine/Prague/1/56
(Sovinova, O., et al, Acta Virol (English ed.) (1958)
2:52-61) (designated herein EIV-A1) and
A/equine/Miami/1/63 (Waddell, G.H., et al, J Am Vet Med
30 Assoc (1963) 143:587-590) (designated herein EIV-A2).
These strains share the characteristics associated with
influenza viruses in general, including those viruses
responsible for human influenza. Most importantly, the
immunological characteristics of influenza virions

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appear to reside primarily in two virally encoded glycoproteins, the hemagglutinin protein (HA) and the neuraminidase protein (NA), both of which are embedded in the membranous envelope which comprises the outer layer of the virus. These proteins attach themselves to the outer membrane of infected cells. Twelve subtypes of HA (H1-H12) and nine subtypes of NA (N1-N9) have been defined using serological cross-reactivity. Thus, all influenza virus isolates carry a parenthetical description (HxNy) corresponding to the subclasses carried. The two equine strains are designated H7N7 (for EIV-A1) and H3N8 (for EIV-A2). (The designation H3 now is used to include both human H3 and former equine HEq2, although these are not identical proteins. H7 is now used to include both avian Hav7 and former HEq1; these, too, are not identical. N7 is former NEq1 and N8 is former NEq2. (Melnick, J., Prog Med Virol (1980) 26:214-232).)

Both HA and NA are involved in the disease process. HA functions in attachment to host membrane and penetration into the host's cells; immunization in subject experimental animals with HA alone elicits neutralizing antibodies and protection from the disease. NA functions in cell-to-cell transmission, and antisera raised against NA attenuate the disease and decrease the spread of the disease from cell to cell. (Schulman, J.L., in The Influenza Viruses and Influenza (1975), E.D. Kilbourne, ed, New York/London: Academic Press, pp. 373-393.)

Human influenza virus and the design of vaccines to protect against it have received considerable attention. Synthetic peptides designed to correspond to a putative antigenic epitope on human HA glycoproteins have been attached to carrier proteins and



used to immunize mice against infection with the human influenza virus of the same strain (U.S. Patent 4,474,757). The peptides synthesized were apparently designed based on the sequence of fragments generated by CNBr digestion of the hemagglutinin protein. Synthetic peptide-carrier vaccines have also been suggested as a general approach to protection against viral infection (Brown, F., Ann Rev Microbiol (1984) 3:221-235).

An alternative approach to vaccine compositions which has been suggested recently utilizes the vaccinia virus as a carrier. Mackett, M., et al, J Virol (1984) 49:857-864 describes this general method. Briefly, vaccinia is a large (187 kb) double-stranded DNA virus which replicates in the cytoplasm of infected cells. It is noninfectious when deproteinized, as it carries its own enzymes for transcription and cannot utilize the machinery of the host cell for this purpose. Vaccinia virus per se was used as the original smallpox vaccine, and is highly desirable as a vaccine carrier because of its low cost and ease of propagation. Freeze-dried vaccinia virus used against smallpox could be mass produced for as little as two cents per dose, while other subunit vaccines, for example, those against hepatitis B, cost approximately \$100 per course of immunization. There are other advantages as well. Vaccinia stimulates both the humoral antibody and cell-mediated immunity systems of the subject. The freeze-dried vaccine is stable without refrigeration and is generally potent after a single inoculation. It is also easy to administer under nonsterile field conditions (Smith, G.L., et al, Biotechniques (1984) Nov/Dec:306-312). Because of these advantages, vaccinia has been used as a carrier for antigenic proteins of hepatitis B and herpes simplex (Paoletti, E., et al,

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Proc Natl Acad Sci (USA) (1984) 81:193-197), rabies
(Wiktor, T.J., et al, Proc Natl Acad Sci (USA) (1984)
81:7194-7198), and human influenza hemagglutinin
(Panicali, D., et al, Proc Natl Acad Sci (1983)
5 80:5364-5368; Smith, G.L., et al, Proc Natl Acad Sci
(1983), 80:7155-7159).

One reason these techniques have not been
extensible to equine influenza is the absence of
sufficient information on the HA and NA proteins of
10 these viruses. This deficiency is remedied by the
present invention, which provides complete genomic and
amino acid sequences for the four relevant surface
proteins characterizing EIV. In addition, the invention
provides properly designed vectors and peptides based on
15 these sequences useful in vaccine compositions.

Disclosure of the Invention

The invention provides complete genetic
sequences encoding the four glycoproteins which
20 characterize the two identified serotypes of EIV. The
invention further provides vaccine compositions which
are grounded in this genetic information. One class of
vaccine compositions comprises modified vaccinia
engineered to express the HA and NA surface
25 glycoproteins of the equine infective agents. Another
class of compositions comprises synthetic peptide
sequences designed to correspond to the antigenic
determinants of the HA proteins. These synthetic
peptides can be size enhanced, such as by linkage to
30 carrier proteins, to engender an immune response. Still
another class comprises the HA or NA proteins per se,
which are preferably produced recombinantly in host
cells which permit correct post-translational
processing. The vaccine compositions of the invention

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may be used in appropriate protocols for protection of horses against the disease, either in independent vaccine regimes, or as adjuncts to more traditional immunizations using killed virus vaccines. A variety of
5 protocols are workable, and several optimal methods based on these protocols form a part of the invention.

The genetic information contained in the cloned cDNAs representing the EIV glycoproteins is useful in two other ways--to construct diagnostic probes for
10 detection of the disease and to construct probes to obtain new cDNAs associated with mutated forms of the virus.

Thus, in one aspect the invention is directed to recombinant DNA comprising the sequences encoding
15 equine H7, equine H3, equine N7, and equine N8. These recombinant sequences or substantial portions thereof are useful as diagnostic probes for detecting the presence of the disease and as probes to retrieve corresponding cDNA in mutated forms of the virus. These
20 sequences or immunologically effective portions thereof are useful in preparing vaccines against EIV which are constructed by inserting them into nonessential portions of the vaccinia virus genome. Accordingly, in another aspect, the invention relates to the resulting
25 vaccinia-borne compositions. In still another aspect, the invention relates to specific synthetic peptide sequences useful in preparing vaccines effective against EIV. The latter vaccines are prepared either by using these peptides per se or by polymerizing these synthetic
30 sequences or attaching them to carrier proteins to provide enhanced size for immunogenicity. The resulting vaccines also form an aspect of the invention. In still another aspect, the invention relates to vaccines comprised of the HA and/or NA proteins per se. In

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addition, the invention includes methods to immunize horses against EIV infection by administration of the foregoing compositions, and to prepare vaccines against mutated forms of the virus.

5

Brief Description of the Drawings

Figures 1-4 show the coding sequences and deduced amino-acid sequences representing the genes and proteins corresponding to H7 (EIV-A1), H3 (EIV-A2), N7 (EIV-A1), and N8 (EIV-A2), respectively.

10

Figure 5 shows the general method to prepare recombinant vaccinia capable of expressing the EIV derived genes.

15

Figure 6 shows the construction of vectors for preparing recombinant vaccinia bearing the coding sequences of Figures 1-4.

Modes of Carrying Out the Invention

A. Definitions and Description of Surface Viral Proteins

20

Hemagglutinin (HA) in general is a glycoprotein of molecular weight approximately 77 kD, and can be cleaved by protease action and subsequent reduction of the single disulfide bond into an amino terminal HA1 50 kD portion and the carboxy terminal HA2 27 kD portion. Complete nucleotide sequences have been determined for at least four avian and human subtypes: H1, H2, H3, and H7. The primary amino acid sequences of all these HA proteins contain highly conserved cysteine, proline, and glycine residues and glycosylation sites.

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Crystallographic studies show that human H3 contains a globular head which extends away from the cellular membrane composed almost entirely of residues of HA1, and an anchoring stalk portion which is comprised of mostly HA2 (Wilson, et al, Nature (1981)

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289:366-373). This is believed to be representative for all HA molecules. Four major antigenic sites, all located in the HA1 globular portion, have been identified from less conserved sequences in immunological variants (Wiley, et al, Nature (1981) 289:373-378). As will be seen below, the amino acid sequence of the equine HA polypeptide differs from those previously determined for the forms infective in humans, particularly in the area of the antigenic sites. In addition, HA peptides in general appear to be subject to antigenic drift with time, although the HA protein from EIV-A1 appears relatively stable.

Neuraminidase (NA) is an approximately 56 kD glycoprotein which occurs as a tetramer on the surface of infected cells. Unlike HA, the membrane anchor is at the amino terminus, and the reactive antigenic sites are predominantly in the carboxy terminal two-thirds of the peptide. Comparison of amino acid sequences from representative N1 and N2 human infective serotypes show extensive homology in the antigenic portion of the molecule, but little in the stalk portion. A similar relationship exists between the EIV sequences shown below.

The equine HA and NA DNA and polypeptide sequences described below and in the figures are representative of their respective equine subtypes. It is recognized that minor mutational changes may occur without destroying the functionality of the DNA sequences in their function as vaccines per se, or that of the proteins generated by them. It is not intended that equine H3, equine H7, equine N7, and equine N8 be limited to the precise nucleotide and amino acid sequences shown. To fall within the designations, a particular sequence must only be functionally

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substantially identical to those depicted in the figures.

One obvious source of variation which has minimal effect on functionality as a coding sequence resides in the redundancy of the genetic code. Of course, the designations are intended to include variations in the sequence which result from this codon redundancy. In addition, minor modifications of the amino acid sequence, and thus the coding sequence, may be permitted which still result in proteins capable of the immunological effect desired. Such changes may include deletion, addition, or alteration of a limited number of amino acids. In particular, for use as a probe, only a substantial portion, and not all of the DNA sequence, is required to be effective. Similarly, for use in a vaccinia-borne vaccine, only an "immunologically effective" portion--i.e., sufficient to generate a protein capable of raising neutralizing antibodies--is required. So long as the altered amino acid sequence functions immunologically in the same fashion as the corresponding sequence depicted, it is included in the designation.

It should be understood that "derived from", when referring to a DNA sequence or amino acid sequence, indicates a correspondence in composition to the referenced material, and not necessarily actual physical derivation therefrom. For example, a DNA encoding a signal sequence which is derived from influenza hemagglutinin refers to a DNA constructed so as to encode a peptide sequence substantially similar to that which is found in influenza HA, and that DNA sequence may be constructed by, for example, obtaining cDNA from the mRNA encoding the protein, by synthetic methods using automated oligonucleotide synthesis, or in any

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other manner designed to recreate a coding sequence for the desired peptide.

B. General Method

5 Based on the genomic information provided by the invention, three general approaches to preparing vaccines are used: in one, partial or full-length HA and NA cDNA clones are recombined into a nonessential region of vaccinia virus, which then express the
10 inserted EIV antigen coding sequences. These vaccinia serve as complete EIV vaccines. In a second approach, synthetic peptides designed from deduced amino acid sequences of equine HA, if necessary made immunogenic by size enhancement, are used to prepare vaccines. These
15 vaccines are used to boost the protective immune response from animals previously given a single dose of killed whole virus vaccine. Alternatively, the synthetic peptide vaccines may be given as a primer prior to the inoculation with whole virus. In a third
20 approach, full-length, properly processed HA or NA proteins are used as vaccines.

HA and NA are used as the basis for active components of the claimed vaccines because in the case of humans, it is believed that immunity is conferred by
25 response to these corresponding surface proteins (Couch, R.B., et al, Ann Rev Microbiol (1983) 527-549). In addition, in horses, antisera raised against whole EIV show high titers of antibody specific against HA of the same serotype strain.

30

B.1. Cloning and Sequencing of the EIV Genes

It is known that the genomes of influenza virus consist of eight segments of single-stranded RNA, each of which encodes a particular viral protein. The eight

segments of RNA from each EIV strain were separated by electrophoresis, and the RNA which encodes HA and that which encodes NA were isolated. cDNA libraries prepared from total viral RNA by conventional methods were
5 screened using reverse transcripts of these RNA isolates and influenza-specific synthetic oligonucleotides as probes. The cDNA clones thus obtained are useful in future preparation of corresponding cDNA for use in vaccines from EIV isolates which have been altered by
10 genetic drift. They thus provide initial probes which enable the production of a series of vaccines corresponding to the inevitable evolutionary drift of the virus.

The positively hybridizing cDNA clones were
15 then sequenced and their identity as HA and NA genes was confirmed by the similarity of the deduced amino acid sequence to known HA and NA proteins. The deduced amino acid sequences were analyzed for antigen sites using secondary structure computer profiles, and by localizing
20 sites known to be important in the disease process, e.g., receptor binding sites.

The resulting cDNAs encoding HA and NA are useful as diagnostic probes for the detection of influenza in horses, as well as in formulating
25 vaccinia-borne vaccines. The probes are used to detect the EIV genome in biological samples such as blood or urine by their ability to hybridize to the target polynucleotide under stringent conditions.

30 B.2. Generation of Recombinant EIV-Vaccinia Viruses

The parameters for effecting the integration of foreign DNA into vaccinia are known in the art and are currently in a position for practical utilization.

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Briefly, the desired immunogen is transferred into a nonessential portion of the vaccinia genome by coinfecting host cells with both native vaccinia and a carrier plasmid which contains the foreign gene sandwiched between sequences homologous with the selected nonessential portion of the vaccinia genome. In addition, the foreign gene is provided with a vaccinia promoter, which will permit its expression under the influence of the vaccinia transcription and translation systems. A general purpose vector capable of housing the foreign gene has been disclosed by Moss, B., et al (Proc Natl Acad Sci (USA) (1983) 80:7155-7159). A diagram of this vector, pGS20, and its manner of use to obtain a recombinant vaccinia virus containing a desired foreign gene is shown in Figure 5.

pGS20 has a vector fragment derived from pBR328, a vector compatible with E. coli which contains the E. coli origin of replication and the ampicillin resistance gene. The vector contains the promoter from the 7.5 k gene of vaccinia (a gene encoding a 7500 dalton protein), excised as a 275 bp HincII/RsaI fragment (Venkatsen, et al, Cell (1981) 25:805-813), which is translocated into the EcoRI site of the vaccinia thymidine kinase (TK) gene. Other restriction site modifications have been made for convenience, and there are BamHI and SmaI restriction sites immediately downstream of the promoter to permit foreign gene cloning (Mackett, M., et al, supra, incorporated herein by reference). Figure 6 shows the nucleotide sequence in the region of the promoter/restriction site fragment junction.

In preparing the vectors of the invention, the DNA encoding the desired immunogen is inserted into pGS20 using the restriction sites downstream of the

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promoter. In the present invention this DNA is derived from equine H7, H3, N7, or N8 and comprises an immunologically effective portion thereof. The recombination vector is amplified in E. coli using transformation to Amp^R and then coinfectd along with wild-type vaccinia into CV-1 cells. Unlike other large DNA animal viruses, vaccinia transcribes and replicates its genome in the cytoplasm of infected cells. Many of the enzymes involved in its nucleic acid metabolism, such as DNA and RNA polymerases, enzymes to cap, methylate, and polyadenylate RNA, as well as thymidine kinase, are encoded in its own genome. Indeed, protein-free vaccinia is noninfective--since it encodes its own transcriptase and apparently cannot use the transcriptase used by its eukaryotic host, it cannot synthesize required proteins using its DNA alone in combination with the host cell machinery. Cells having been transformed with both the recombinant, for example pGS20-derived, vector and vaccinia virus mediate recombination of the DNA portion contained within the vaccinia-corresponding segments of the vector into the vaccinia genome. Use of the TK-encoding sequences to effect recombination is particularly desirable, as not only is this a nonessential portion of the vaccinia genome, but a selectable marker is provided for cells containing the recombined vaccinia--i.e., when the foreign DNA is inserted, the TK gene is inactivated, and the tk- recombinant viruses can be selected by plaque assay on tk- cells in the presence of 5-bromodeoxyuridine (BUdR). As shown in Figure 5, tk-recombinant virus plaques can be selected yielding the desired recombinant vaccinia. Additionally, tk-vaccinia have been shown to be 10^5 to 10^6 times less virulent in animals as compared to wild-type vaccinia.

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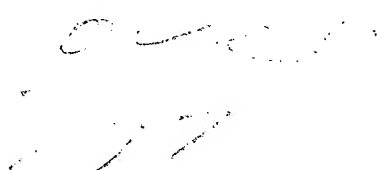
indicating the potential for an increased safety factor in humans (Smith, G.L., and Moss, B., Biotechniques (1984) 306-312).

While pGS20 is a convenient illustrative
5 vector, it is understood that alternative constructions involving other vaccinia promoters in other nonessential regions of the gene may be used (Moss, et al, Gene Amplification Analysis, Vol. III, Pappas, T.K., et al, eds. (1982) New York, Elsevier, pp. 201-213; Mackett,
10 M., et al, supra). Not only CV-1, but any competent cells susceptible to vaccinia infection can be used to effect the recombination.

B.3. Preparation of Synthetic Peptides

15 Peptides were identified as representing antigenic regions of the HA viral proteins by analogy to the corresponding regions in the human influenza proteins, by analysis of the predicted secondary structure of the equine proteins, and by location of key
20 functional areas in the equine proteins. These potential antigenic determinants were synthesized using the solid-state Merrifield method. As these peptides are often too small to be immunogenic, their size is enhanced either by polymerization or by conjugation to a
25 carrier protein.

A variety of techniques is available in the art for such polymerization or conjugation. The most commonly used polymerization technique employs glutaraldehyde and effects self-condensation of the
30 peptide. However, conjugation to a carrier protein is, generally, a preferred approach. Such conjugation is effected by linking agents which attach to each member of the conjugate. For example, there are a large number of hetero bifunctional agents which generate a disulfide



link with one component and a peptide link with the other. These have been extensively used. Any of the synthetic peptides disclosed herein may be prepared with an additional cysteine to permit this mode of linkage.

5 The most popular such linking agent is N-succinidyl-3-(2-pyridylthio) propionate (SPDP). Other such agents, which create a disulfide linkage at one end and an amide linkage through an ϵ amino on a lysine at the other end, are available. See, for example, Immun
10 Rev (1982) 82:185. Still other bifunctional coupling agents form a thioether rather than a disulfide linkage. Many of these thioether-forming agents are commercially available, and include reactive esters of 6-maleimidocaproic acid, 2-bromoacetic acid,
15 2-iodoacetic acid, and the like. Carboxyl groups of these acids can be activated by combining them with succinimide or 1-hydroxy-2-nitro-4-sulfonic acid, sodium salt. In addition, reagents which operate through dehydration, such as dicyclohexylcarbodiimide, can be
20 used.

B.4. Preparation of Recombinant HA or NA Vaccines

In addition to the vaccines prepared by
25 obtaining recombinant vaccinia virus and by size enhancement of synthetic peptides, larger proteins derived from the HA or NA proteins of the equine influenza viral coat proteins can be prepared. As the entire genome encoding these proteins is available,
30 suitable DNA encoding these proteins in their entirety or such smaller portions thereof which are immunologically effective may be excised from cDNA and ligated into expression vectors for production of these proteins in foreign hosts. While it is conceivable that

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bacterial hosts can be used, the production of these proteins in mammalian cells lines, or even yeast cells, is preferred, since post-translational processing available in these hosts more accurately simulates that ordinarily carried out in cells infected with the virus. Therefore the recombinant proteins produced would be expected to more closely mimic the viral proteins as ordinarily found by a target cell.

Suitable host mammalian and yeast cells, and appropriate control sequences and transfer vectors effective in such hosts, are found in C.5 below. In general, the desired portion of the protein-encoding DNA sequence is excised from the gene using suitable restriction enzymes, and modified, if necessary, prior to ligating into transfer vectors providing operable control sequences. The ligated vectors are then transformed into the corresponding hosts, and the hosts cultured under conditions known to foster the multiplication of the host cell, and, usually, followed by inducing conditions to activate the control sequences provided. The recombinant HA and NA proteins are then retrieved from the cells, by lysis, if necessary, and purified using standard techniques to obtain the pure recombinant protein.

25 In addition, the desired recombinant protein may be produced in cell culture using co-infection with vaccinia virus along with the appropriate recombinant pGS20 derivative (Cochran, et al, Proc Natl Acad Sci (USA) (1985) 82:19-23).

B.5. Vaccine Preparation

The vaccines of the invention which are recombined vaccinia virus are employed in the conventional manner. Vaccinia virus can be cultured in

standard media, as described below, or by infecting live animals and harvesting the virus from the lesions. The administration of the vaccinia is generally by means of scratching the skin, and non-sterile conditions are often acceptable.

For vaccines which contain peptides or proteins as active ingredients, administration is slightly more complex. Typically, such vaccines are prepared as injectables, either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid prior to injection may also be prepared. The preparation may also be emulsified. The active immunogenic ingredient is often mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredient. Suitable excipients are, for example, water, saline, dextrose, glycerol, ethanol, or the like, and combinations thereof. In addition, if desired, the vaccine may contain minor amounts of auxiliary substances such as wetting or emulsifying agents, pH buffering agents, or adjuvants which enhance the effectiveness of the vaccine. The vaccines are conventionally administered parenterally, by injection, for example, either subcutaneously or intramuscularly. Additional formulations which are suitable for other modes of administration include suppositories, intranasal aerosols, and, in some cases, oral formulations. For suppositories, traditional binders and carriers may include, for example, polyalkylene glycols or triglycerides; such suppositories may be formed from mixtures containing the active ingredient in the range of 0.5% to 10%, preferably 1%-2%. Oral formulations include such normally employed excipients as, for example, pharmaceutical grades of manitol, lactose, starch, magnesium stearate, sodium saccharin,

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cellulose, magnesium carbonate, and the like. These compositions take the form of solutions, suspensions, tablets, pills, capsules, sustained release formulations, or powders, and contain 10%-95% of active ingredient, preferably 25%-70%.

The peptides' proteins may be formulated into the vaccine as neutral or salt forms. Pharmaceutically acceptable salts include the acid addition salts (formed with the free amino groups) and which are formed with inorganic acids such as, for example, hydrochloric or phosphoric acids, or such organic acids as acetic, oxalic, tartaric, mandelic, and the like. Salts formed with the free carboxyl groups may be derived from from inorganic bases such as, for example, sodium, potassium, ammonium, calcium, or ferric hydroxides, and such organic bases as isopropylamine, trimethylamine, 2-ethylamino ethanol, histidine, procaine, and the like.

The peptide or protein-based vaccines are administered in a manner compatible with the dosage formulation, and in such amount as will be therapeutically effective and immunogenic. The quantity to be administered depends on the subject horse to be treated, capacity of the subject's immune system to synthesize antibodies, and the degree of protection desired. Precise amounts of active ingredient required to be administered depend on the judgment of the practitioner and are peculiar to each subject. However, suitable dosage ranges are of the order of several hundred micrograms active ingredient per subject. Suitable regimes for initial administration and booster shots are also variable, but are typified by an initial administration followed in one or two week intervals by a subsequent injection or other administration. Alternatively, primer doses of the vaccines of the

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invention may be used one to two weeks before injection with conventional killed virus vaccine.

Administration is typically by injection intramuscularly, but the peptides or proteins may also
5 be formulated for intranasal application.

A preferred use of the synthetic peptide immunogens is that they be incorporated into existing immunization regimens as either a first or "priming" inoculation or as second and subsequent "boosting"
10 inoculations in combination with other vaccines. It has been shown by others that a single dose of a synthetic peptide "primes" the immune system such that subsequent inoculation with the whole virus (e.g., poliovirus) elicits extremely high titers of neutralizing
15 antibodies. The practice of priming or boosting with synthetic peptides offers two major advantages: it lessens the number of whole virus vaccine inoculations (and the accompanying side effects) required to achieve immunity, and it retains at least one dose of whole
20 virus vaccine in the immunization regimen, thus allowing the animal's immune system an opportunity to see the entire antigenic repertoire of the pathogen.

C. Standard Methods

25 C.1. Preparation of cDNA

First-strand cDNA was synthesized using RNA-dependent DNA polymerase from avian myeloblastosis virus (AMV), according to Ulrich, A., et al, Science (1977) 196:1313-1319, using an appropriate primer. RNA
30 template was removed by denaturation at 100°C for five minutes followed by chilling on ice and centrifugation at 5000 rpm for 10 min at room temperature. The supernatant was treated with DNA polymerase I, large fragment to synthesize the second strand. The hairpin

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loop was cleaved with S1 nuclease from Aspergillus
oryzae in 300 mM NaCl, 30 mM NaOAc, pH 4.5, 3 mM ZnCl₂
for 30 min at 37°C with 600 units of enzyme. After
extraction of cDNA with phenol/chloroform, small
5 oligonucleotides were removed by three ethanol
precipitations in the presence of ammonium acetate,
i.e., using 1/2 volume 7.5 M ammonium acetate and two
volumes of ethanol at -70°C. The resulting blunt-ended,
double-stranded cDNA can then be fractionated either
10 using gel filtration through a Sepharose 4B (Pharmacia)
or by ultracentrifugation in 5-20% gradients of glycerol
in 100 mM NaCl, 10 mM Tris-HCl, pH 7.6, and 1 M EDTA,
followed by fractionation of the gradient.

The double-stranded DNA was then cloned into
15 suitable vectors using homopolymeric tailing essentially
as described by Sutcliffe, J.G., Nucleic Acid Res (1978)
5:2721-2732.

Thus, to complete cDNA libraries in pBR322,
double-stranded cDNAs were tailed with dCTP using a
20 reaction containing 0.2 M potassium cacodylate, 25 mM
Tris, pH 6.9, 2 mM dithiothreitol, 0.5 mM CoCl₂, 200
mM dCTP, 400 µ/ml BSA, and 40 units calf thymus
terminal transferase for 5 min at 22°C. After
extraction with phenol/chloroform and ethanol
25 precipitation the dC-tailed cDNA was annealed with
PstI-digested pBR322 tailed with oligo-dG using 2.5 µg
pBR322 per 1 µg cDNA/ml. The annealed mixture was
transferred into E. coli 1061 using the CaCl₂
treatment of Casabadian, N., et al, J Mol Biol (1980)
30 138:179-207.

C.2. Screening cDNA Libraries

cDNA libraries were screened by replicating
plates onto duplicate nitrocellulose filters (S&S BA-85)

and permitting the colonies to grow at 37°C for 14-16 hr on L-agar containing 15 µg/ml tetracycline, followed by 12-24 hr on L-agar containing 170 µg/ml chloramphenicol. Colonies were lysed with 10% SDS, and
5 the DNA fixed to the filter by sequential treatment for 5 min with 0.5 M NaOH, 1.5 M NaCl, then 0.5 Tris-HCl, pH 8.0, 1.5 M NaCl, followed by 2x standard saline citrate (1x SSC = 0.5 M NaCl, 0.105 M sodium citrate). Filters were air-dried and baked at 80°C for 2 hr in vacuo.

10 For nick-translated probe, the duplicate filters are prehybridized at 42°C for 16-18 hr with 10 ml per filter of DNA hybridization buffer (50% formamide (40% formamide if reduced stringency), 5x SSC, pH 7.0, 5x Denhardt's solution (polyvinylpyrrolidone, plus
15 Ficoll and bovine serum albumin; 1x = 0.02% of each), 50 mM sodium phosphate buffer at pH 7.0, 0.2% SDS, 50 µg/ml yeast tRNA, and 50 µg/ml denatured and sheared salmon sperm DNA).

Samples are hybridized with nick-translated DNA
20 probes at 42°C for 12-36 hr for homologous species and 37°C for heterologous species contained in 5 ml of this same DNA hybridization buffer. The filters are washed two times for 30 min, each time at 50°C, in 0.2x SSC, 0.1% SDS for homologous species hybridization, and at
25 50°C in 3x SSC, 0.1% SDS for heterologous species hybridization. Filters are air dried and autoradiographed for 1-3 days at -70°C.

For synthetic (12-30 mer) oligonucleotide probes, the duplicate filters are prehybridized at 42°C
30 for 2-8 hr with 10 ml per filter of oligo-hybridization buffer (6x SSC, 0.1% SDS, 1 mM EDTA, 5x Denhardt's, 0.05% sodium pyrophosphate and 50 µg/ml denatured and sheared salmon sperm DNA).

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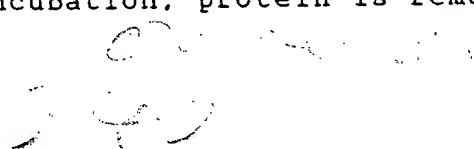
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The samples are hybridized with kinased oligonucleotide probes of 12-30 nucleotides under conditions which depend on the composition of the oligonucleotide. Typical conditions employ a
5 temperature of 30-42°C for 24-36 hr with 5 ml/filter of this same oligo-hybridization buffer containing probe. The filters are washed two times for 15 min at 23°C, each time with 6x SSC, 0.1% SDS and 50 mM sodium
10 phosphate buffer at pH 7, then are washed once for 2 min at the calculated hybridization temperature with 6x SSC and 0.1% SDS, air dried, and are autoradiographed at -70°C for 2 to 3 days.

C.3. Vector Construction and Verification

15 Construction of suitable vectors containing the desired coding and control sequences employs standard ligation and restriction techniques which are well understood in the art. Isolated plasmids, DNA
sequences, or synthesized oligonucleotides are cleaved,
20 tailored, and religated in the form desired.

Site specific DNA cleavage is performed by treating with the suitable restriction enzyme (or
enzymes) under conditions which are generally understood in the art, and the particulars of which are specified
25 by the manufacturer of these commercially available restriction enzymes. See, e.g., New England Biolabs, Product Catalog. In general, about 1 µg of plasmid or DNA sequence is cleaved by one unit of enzyme in about
20 µl of buffer solution; in the examples herein,
30 typically, an excess of restriction enzyme is used to insure complete digestion of the DNA substrate. Incubation times of about one hour to two hours at about 37°C are workable, although variations can be tolerated. After each incubation, protein is removed by



extraction with phenol/chloroform, and may be followed by ether extraction, and the nucleic acid recovered from aqueous fractions by precipitation with ethanol. If desired, size separation of the cleaved fragments may be performed by polyacrylamide gel or agarose gel electrophoresis using standard techniques. A general description of size separations is found in Methods in Enzymology (1980) 65:499-560.

Restriction cleaved fragments may be blunt ended by treating with the large fragment of E. coli DNA polymerase I (Klenow) in the presence of the four deoxynucleotide triphosphates (dNTPs) using incubation times of about 15 to 25 min at 20 to 25°C in 50 mM Tris pH 7.6, 50 mM NaCl, 6 mM $MgCl_2$, 6 mM DTT and 5-10 μM dNTPs. The Klenow fragment fills in at 5' sticky ends but chews back protruding 3' single strands, even though the four dNTPs are present. If desired, selective repair can be performed by supplying only one of the, or selected, dNTPs within the limitations dictated by the nature of the sticky ends. After treatment with Klenow, the mixture is extracted with phenol/chloroform and ethanol precipitated. Treatment under appropriate conditions with S1 nuclease or Bal-31 results in hydrolysis of any single-stranded portion.

Synthetic oligonucleotides are prepared by the method of Efimov, V.A., et al (Nucleic Acids Res (1982) 6875-6894), and can be prepared using commercially available automated oligonucleotide synthesizers. Kinasing of single strands prior to annealing or for labeling is achieved using an excess, e.g., approximately 10 units of polynucleotide kinase to 1 nmole substrate in the presence of 50 mM Tris, pH 7.6, 10 mM $MgCl_2$, 5 mM dithiothreitol, 1-2 mM ATP, 1.7

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pmoles $\gamma^{32}\text{P}$ -ATP (2.9 mCi/mmole), 0.1 mM spermidine, 0.1 mM EDTA.

Ligations are performed in 15-50 μl volumes under the following standard conditions and temperatures: 20 mM Tris-Cl pH 7.5, 10 mM MgCl_2 , 10 mM DTT, 33 $\mu\text{g/ml}$ BSA, 10 mM-50 mM NaCl, and either 40 μM ATP, 0.01-0.02 (Weiss) units T4 DNA ligase at 0°C (for "sticky end" ligation) or 1 mM ATP, 0.3-0.6 (Weiss) units T4 DNA ligase at 14°C (for "blunt end" ligation). Intermolecular "sticky end" ligations are usually performed at 33-100 $\mu\text{g/ml}$ total DNA concentrations (5-100 nM total end concentration). Intermolecular blunt end ligations (usually employing a 10-30 fold molar excess of linkers) are performed at 1 μM total ends concentration.

In vector construction employing "vector fragments", the vector fragment is commonly treated with bacterial alkaline phosphatase (BAP) or calf intestinal alkaline phosphatase (CIP) in order to remove the 5' phosphate and prevent religation of the vector. Digestions are conducted at pH 8 in approximately 150 mM Tris, in the presence of Na^+ and Mg^{+2} using about 1 unit of BAP or CIP per μg of vector at 60° for about one hour. In order to recover the nucleic acid fragments, the preparation is extracted with phenol/chloroform and ethanol precipitated. Alternatively, religation can be prevented in vectors which have been double digested by additional restriction enzyme digestion of the unwanted fragments.

Ligations for plasmid construction are confirmed by first transforming E. coli strain MC1061 obtained from Dr. M. Casadaban (Casadaban, M., et al, J Mol Biol (1980) 138:179-207) or other suitable host with the ligation mixture. Successful transformants are

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selected by ampicillin, tetracycline or other antibiotic resistance or using other markers depending on the mode of plasmid construction, as is understood in the art. Plasmids from the transformants are then prepared according to the method of Clewell, D. B., et al, Proc Natl Acad Sci (USA) (1969) 62:1159, optionally following chloramphenicol amplification (Clewell, D. B., J Bacteriol (1972) 110:667). The isolated DNA is analyzed by restriction and/or sequenced by the dideoxy method of Sanger, F., et al, Proc Natl Acad Sci (USA) (1977) 74:5463 as further described by Messing, et al, Nucleic Acids Res (1981) 9:309, or by the method of Maxam, et al, Methods in Enzymology (1980) 65:499.

15 C.4 Site-Specific Mutagenesis

For portions of vectors derived from cDNA or genomic DNA which require sequence modifications, site specific primer directed mutagenesis is used. This is conducted using a primer synthetic oligonucleotide complementary to a single stranded phage DNA to be mutagenized except for limited mismatching, representing the desired mutation. Briefly, the synthetic oligonucleotide is used as a primer to direct synthesis of a strand complementary to the phage, and the resulting double-stranded DNA is transformed into a phage-supporting host bacterium. Cultures of the transformed bacteria are plated in top agar, permitting plaque formation from single cells which harbor the phage.

30 Theoretically, 50% of the new plaques will contain the phage having, as a single strand, the mutated form; 50% will have the original sequence. The resulting plaques are hybridized with kinased synthetic primer and then washed at a temperature which permits

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hybrids of an exact match to remain, but at which the mismatches with the original strand are washed off. Plaques which remain hybridized to the probe at the stringent wash temperature are then picked, cultured, and the DNA recovered.

For probing, plaques are screened by replicating the plaques onto duplicate nitrocellulose filter papers (S & S type BA-85) and infected cells are allowed to grow at 37°C for 14-16 hr on L agar containing 15 µg/ml tetracycline. The colonies are lysed with 10% SDS and the DNA is fixed to the filter by sequential treatment for 5 min with 500 mM NaOH/1.5 M NaCl, then 0.5 M Tris HCl(pH 8.0)/1.5 M NaCl followed by 2x standard saline citrate (SSC). Filters are air dried and baked at 80°C for 2 hr.

For synthetic (12-30 mer) oligonucleotide probes, the duplicate filters are prehybridized at 42°C for 2-8 hr with 10 ml per filter of oligo-hybridization buffer (6x SSC, 0.1% SDS, 1 mM EDTA, 5x Denhardt's, 0.05% sodium pyrophosphate and 50 µg/ml denatured and sheared salmon sperm DNA).

The samples are hybridized with kinased oligonucleotide probes of 12-30 nucleotides under conditions which depend on the composition of the oligonucleotide. Typical conditions employ a temperature of 30-42°C for 24-36 hr with 5 ml/filter of this same oligo-hybridization buffer containing probe. The filters are washed two times for 15 min at 23°C, each time with 6x SSC, 0.1% SDS and 50 mM sodium phosphate buffer at pH 7, then are washed once for 2 min at the stringent wash temperature with 6x SSC and 0.1% SDS. Typically, the stringent wash temperature for oligonucleotides of 16-24 bases with from 1 to 3 mismatches will be 40-70°C, and can most easily be

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determined by successive washes of the hybridized filter. For example, the hybridized filters can be washed first at 40°C, then at 50°C, then at 60°C, and then at 70°C, with air drying of the filter and
5 autoradiography at -70°C overnight between each wash.

A shorter modification of this procedure was particularly useful in the construction of mutated sequences containing new restriction sites. In this modification, the sequence to be modified is excised and
10 inserted into M13 vectors as a single strand and the preparation is treated with Klenow in the presence of primer to create a heteroduplex, as in the above conventional procedure. The insert is then removed from the phage vector and cloned into a convenient host
15 plasmid, such as a pUC or pBR plasmid for amplification. After amplification the plasmid DNA is cut with a restriction enzyme protocol which includes the enzyme recognizing the desired new site. Only the mutagenized double-stranded plasmids yield the correct
20 length fragment. These fragments are isolated, for example on gels, and no probing is necessary.

C.5. Hosts and Control Sequences

Both procaryotic and eucaryotic systems may be
25 used to express the EIV encoding sequences; procaryotic hosts are the most convenient for cloning procedures. Procaryotes most frequently are represented by various strains of E. coli; however, other microbial strains may also be used. Plasmid vectors which contain replication
30 sites and control sequences derived from a species compatible with the host are used; for example, E. coli is typically transformed using derivatives of the pUC series or pBR322. pBR322, for example, contains genes for ampicillin and tetracycline resistance, and thus

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provides additional markers which can be either retained or destroyed in constructing the desired vector.

Commonly used procaryotic control sequences which are defined herein to include promoters for transcription
5 initiation, optionally with an operator, along with ribosome binding site sequences, include such commonly used promoters as the beta-lactamase (penicillinase) and lactose (lac) promoter systems (Chang, et al, Nature
10 (1977) 198:1056 and the tryptophan (trp) promoter system (Goeddel, et al Nucleic Acids Res (1980) 8:4057 and hybrid control systems such as those employing upstream trp and downstream lac regions.

Eucaryotic hosts are often preferred because of their ability to effect post-translational processing.
15 Microbial eucaryotic hosts include yeast; laboratory strains of Saccharomyces cerevisiae, Baker's yeast, are most used although a number of other strains are commonly available. Vectors employing, for example, the 2 μ origin of replication of Broach, J. R., Meth Enz
20 (1983) 101:307, or other yeast compatible origins of replications (see, for example, Stinchcomb, et al, Nature (1979) 282:39, Tschempe, et al, Gene (1980) 10:157 and Clarke, L. et al, Meth Enz (1983) 101:300) may be used. Control sequences for yeast vectors
25 include promoters for the synthesis of glycolytic enzymes (Hess, et al, J Adv Enzyme Req (1968) 7:149; Holland, et al, Biochemistry (1978) 17:4900). Additional promoters known in the art include the promoter for 3-phosphoglycerate kinase (Hitzeman, et al,
30 J Biol Chem (1980) 255:2073), and those for other glycolytic enzymes.

Eucaryotic host cell cultures derived from multicellular organisms may be used also. These systems have the additional advantage of the ability to splice

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out introns and thus can be used directly to express genomic fragments. Useful host cell lines include VERO and HeLa cells, and Chinese hamster ovary (CHO) cells. Expression vectors for such cells ordinarily include
5 promoters and control sequences compatible with mammalian cells such as, for example, the commonly used early and late promoters from Simian Virus 40 (SV 40) (Fiers, et al, Nature (1978) 273:113), or other viral
10 promoters such as those derived from polyoma, Adenovirus 2, bovine papiloma virus, or avian sarcoma viruses. The controllable promoter, hMTII (Karin, M., et al, Nature (1982) 299:797-802) may also be used. General aspects of mammalian cell host system transformations have been described by Axel; U.S. Patent No. 4,399,216 issued 16
15 August 1983. It now appears, also that "enhancer" regions are important in optimizing expression; these are, generally, sequences found upstream or downstream of the promoter region in non-coding DNA regions. Origins of replication may be obtained, if needed, from
20 viral sources. However, integration into the chromosome is a common mechanism for DNA replication in eucaryotes.

Depending on the host cell used, transformation is done using standard techniques appropriate to such
25 cells. The calcium treatment employing calcium chloride, as described by Cohen, S. N., Proc Natl Acad Sci (USA) (1972) 69:2110, or the $RbCl_2$ method described in Maniatis, et al, Molecular Cloning: A Laboratory Manual (1982) Cold Spring Harbor Press, p.
30 254 may be used for procaryotes or other cells which contain substantial cell wall barriers. For mammalian cells without such cell walls, the calcium phosphate precipitation method of Graham and van der Eb, Virology (1978) 52:546, optionally as modified by Wigler, M., et

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al. Cell (1979) 16:777-785 may be used. Transformations into yeast may be carried out according to the method of Van Solingen, P., et al. J Bact (1977) 130:946 or of Hsiao, C. L., et al. Proc Natl Acad Sci (USA) (1979) 76:3829.

C.6. ELISA Assay for Antibodies Against Test Peptides

The synthetic peptides of the invention and their size-enhanced derivatives were used to immunize rabbits and mice, and their ability to raise antibodies in sera was evaluated using an enzyme-linked immunosorbent assay (ELISA). The appropriate peptide for reaction with the antisera was bound to 96-well plates by pretreating the wells in 2% glutaraldehyde for 2 hr at room temperature and then incubating a solution of the appropriate peptide (100 μ l of 50 μ g peptide/ml PBS) in the wells for 2 hr at 37°C. After washing, the wells received fourfold dilutions of antisera ranging from 1/20 to 1/327,680, and the plates were incubated for 2 hr at 37°C. After washing, the wells received a 1/3500 dilution of horseradish peroxidase-conjugated goat anti-rabbit immunoglobulin (Boehringer-Mannheim) in 0.1% ovalbumin and 0.05% Tween-20 in PBS. The plates were incubated at 4°C overnight. After washing, a solution of 0.1% citric acid, pH 5.0, 0.04% o-phenyldiamine, and 0.012% peroxide was added to each well, and the plates were incubated for 5-30 min at room temperature. The color reaction was stopped by adding 4 N HCl and the absorbance at 490 nm was read on a Biotek ELISA Reader, Model EL-308. Titers for each serum were determined by plotting absorbance against serum dilution for both preimmune and immune samples.

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C.7 Plaque Reduction Assay

The ability of antisera to neutralize virus was tested using a plaque reduction assay. Preimmune sera were used as controls. Antisera to be tested and control antisera were heat-treated at 56°C for 30 min in order to inactivate nonspecific antiviral agents present in all mammalian sera. 100-150 plaque-forming units (pfu) of EIV were incubated with an equal volume of appropriately diluted control and test sera (usually twofold dilutions) for 1 hr at 25°C. Monolayers of MCDK cells were then inoculated with the virus/serum mixture and adsorption allowed to take place for 1 hr at room temperature. The monolayers were then rinsed with Earl's basic salt solution and overlaid with a mixture of DMEM and 0.8% agarose containing 2.5 µg/ml trypsin. The wells were incubated in a 5% CO₂ atmosphere at 37°C for 4 days in order allow plaque formation by viable virus. Cells were stained with crystal violet and total plaques recorded. The serum neutralization titer was determined as the dilution of the serum that reduced EIV pfu by greater than 50%.

D. Examples

The following examples are intended to illustrate the invention, but not to limit its scope.

D.1. Preparation and Sequencing of Equine HA and NA cDNA

Both known EIV strains were used as sources of RNA encoding the desired proteins. Any fresh isolate from a diseased animal can be used as starting material; both EIV-A1 and EIV-A2 are pandemic in the diseased equine population. In the examples below, EIV-A1 strain (H7N7) was originally isolated from an infected horse in

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Florida in 1973 and passed in four isolation ponies at Cornell University before being propagated in Madin-Darby canine kidney (MDCK) cells. It is designated A1/Cornell/16/74. The EIV-A2 strain (H3N8), designated A2/Kentucky/1/81, was obtained as an allantoic fluid preparation. Both stocks were shown to produce clinically recognizable disease in horses.

These strains, propagated in MDCK cells, were concentrated by centrifugation and purified by banding twice in 30%/60% discontinuous sucrose gradients. The purified virus was treated with 10 µg/ml DNase at 23°C for twenty minutes, and the virions were disrupted using 0.5% SDS followed by two extractions with phenol:chloroform:isoamyl alcohol (50:50:1) and two extractions with chloroform:isoamyl alcohol (50:1). The aqueous phase was adjusted to 0.3 M sodium acetate, and RNA was precipitated with 2-1/2 volumes absolute ethanol.

Preparation of cDNA Probes

To isolate RNA fragments for cDNA probe preparation, 1.5-2 µg viral RNA was denatured in 10 mM methylmercuric hydroxide for 1.5 minutes at 65°C and then electrophoresed on a 1.4% agarose gel in Tris-borate buffer (0.089 M Tris, 0.089 M boric acid, 0.02 M EDTA) at 12 V/cm for two hours. The gel was stained using 0.5 µg/ml ethidium bromide. The RNA bands identified as HA or NA (by size analogy to human influenza models) were then used to prepare cDNA probes by the method of Lonberg and Gilbert, Proc Natl Acad Sci (USA) (1983) 80:3661-3665. Briefly, 5-10 µl agarose plugs were removed from the appropriate areas of the gel with 50 µl capillary tubes and the plugs melted at 65°C for three minutes and cooled to 37°C. First strand cDNA was obtained using RNA-dependent DNA polymerase, as

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in section C.1. Calf thymus DNA was used as a random primer.

Preparation of cDNA Libraries Containing HA and
NA

5 cDNA libraries in pBR322 were prepared from total RNA of each strain. The cDNA libraries were prepared as described above using two primer methods. In the first method, oligo-dT 12-18mers were used as
10 primers after addition of poly-A tails to the viral negative stranded poly-A deficient RNA. Three µg of viral RNA was incubated with E. coli poly-A polymerase in a reaction containing 50 mM Tris-HCl, pH 7.6, 10 mM MgCl₂, 2.5 mM MnCl₂, 250 mM NaCl, 10 mM ATP for 3
15 min at 37°C and then extracted with phenol/chloroform and ethanol-precipitated. The resulting extended RNA was used as template.

In the alternative method, the synthetic oligomer 5'-AGCAAAAGCAGG-3' (oligo a), which is a known
20 complement to twelve nucleotides in the 3' terminal untranslated region of each of the eight individual viral RNAs in all influenza viruses was used as primer. A second synthetic primer 5'-AGTAGAAACAAGG-3' (oligo b), which complements an influenza consensus sequence in the
25 5' untranslated region at the upstream end of each coding region in the viral RNA, was used to prime the complementary strand. Three µg viral RNA yielded approximately 1 µg cDNA size selected to 500 bp; and each library contained about 500,000 independent
30 recombinants.

For probing with cDNA probes, hybridization buffer contained 50% formamide, 5x SSC, 50 mM HEPES, pH 8.0, 5x Denhardt solution (1x Denhardt's = 0.02% each polyvinyl pyrrolidone, Ficoll, and BSA), 0.1% SDS, 50

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mM/ml yeast tRNA, and 50 mM sheared and denatured salmon sperm DNA. For probing with synthetic probes, the hybridization buffer contained 6x SSC, 0.1% SDS, 1 mM EDTA, 5x Denhardt's, 0.05% sodium pyrophosphate, and 15 μ g/ml denatured and sheared salmon sperm DNA.

Isolation of HA and NA cDNA Sequences

The cDNA libraries were probed using one of the reverse-transcribed RNA for each of the desired H7, H3, N7, and N8 sequences prepared as described above and using the two synthetic oligomers used for priming in the reverse-transcribed synthesis, as also described (oligos a and b). For reverse-transcribed RNA, prehybridization was at 42°C for 12-14 hr, and hybridization was at 42°C for 12-36 hr using 10^5 cpm/filter. For probing with oligos a or b, prehybridization was at 42°C for 2-8 hours and hybridization, at 33°C for 6-12 hours using 10^6 cpm/filter-kinased probe. (The 33°C temperature was calculated using the formula $T_H = 4[G + C] + 2[A + T] - 3^\circ\text{C}$. Suggs, S.V., et al, Developmental Biology Using Purified Genes (1981), ed. D.D. Brown and C.F. Fox, Academic Press, New York, pp. 683-693.) For the RNA probe, the filters were washed twice for 30 min each at 50°C in 0.2x SSC, 0.1% SDS, and then air dried and autoradiographed for 1-3 days at -70°C. For the oligo probes the filters were washed twice for 2 min at 23°C each, with 6x SSC, 0.1% SDS, and 0.05% sodium pyrophosphate, then for 2 min at 33°C in the same buffer, and then air dried and autoradiographed at -70°C for 1-2 days. A large number of positive clones were obtained from both EIV-A1 and EIV-A2 libraries, but only one clone from each library which hybridized with all

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three probes in each of the four cases was retained for further study.

The resulting clones representing H7, H3, N7, and N8 were sequenced by subcloning into M13 mp8, mp9, mp18, and mp19 (Messing, J., et al, Gene (1982) 19:269-276) following sequence determination using the dideoxy method of Sanger, et al, Proc Natl Acad Sci (USA) (1977), 74:5463-5467. The nucleotide sequence of the cDNA sequences for H7, H3, N7, and N8 are shown in Figures 1-4.

Examination of the sequences in Figures 1-4 indicates the characteristic features of HA1 and HA2 proteins from H7 and H3, including conserved cysteines, clip sites between HA1 and HA2, and the conserved membrane fusion region at the amino terminus of HA2.

D.2. Synthesis of Synthetic Peptides Based HA Sequence Information

The following peptides were synthesized using the method of Merrifield, J., J Amer Chem Soc (1963) 85:2149-2154.

Based on the EIV-A1 (H7):

(A1-X) representing residues 146-173: Ala-Cys-Arg-Arg-Ser-Arg-Ser-Ser-Phe-Tyr-Ala-Glu-Met-Lys-Trp-Leu-Leu-Ser-Asn-Thr-Asp-Asn-Gly-Val-Phe-Pro-Gln-Met.

(A1-Y) representing residues 181-209: (Cys)-Lys-Arg-Glu-Pro-Ala-Leu-Ile-Ile-Trp-Gly-Ile-His-His-Ser-Gly-Ser-Thr-Ala-Glu-Gln-Thr-Arg-Leu-Tyr-Gly-Ser-Gly-Asn-Lys-(Cys)

(A1-Z) representing residues 309-337: Arg-Ala-Ile-Gly-Lys-Cys-Pro-Arg-Tyr-Val-Lys-Gln-Lys-Ser-Leu-Met-Leu-Ala-Thr-Gly-Met-Lys-Asn-Val-Pro-Glu-Asn-Ser-Thr.

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For EIV-A2:

(A2-X) representing residues 143-159: Thr-Gly-Val-Thr- Gln-Asn-Gly-Arg-Ser-Gly-Ala-Cys-Arg-Arg-Gly-Ser-Ala-(Ser)-(Arg).

5 (A2-Y) representing residues 191-218: Lys-Leu-Tyr-Ile-Trp-Gly-Ile-His-His-Pro-Ser-Thr-Asn-Asn-Glu-Gln-Thr-Lys-Leu-Tyr-Ile-Gln-Glu-Ser-Gly-Arg-Val-Thr.

(A2-Z) representing residues 322-343: Lys-Tyr-Ile-Lys-Gln-Asn-Thr-Leu-Lys-Leu-Ala-Thr-Gly-Met-Arg-Asn-Val-Pro-Glu-Lys-Gln-Ile.

10 Finally, a synthetic peptide was designed after the amino terminus of the HA2 portion of both the H7 and H3 hemagglutinins (residues 345-357 in H3 and 350-362 in H7):

15 (A1/A2-N) Gly-Ile-Phe-Gly-Ala-Ile-Ala-Gly-Phe-Ile-Glu-Asn-Gly-(Lys)-(Lys).

Residues in parenthesis are not present in natural sequence but were added for purposes of peptide conjugation or ease in synthesis.

20 The foregoing peptides are designated by the shorthand symbols shown, and were selected for synthesis on the following basis: both X-peptides (A1-X and A2-X) and Y-peptides (A1-Y and A2-Y) correspond to regions in the human analogs which are major antigenic sites as
25 mapped by monoclonal antibodies and by sequencing of wild-type variants, and thus represent epitopes which, in the human model, elicit B-cell response.

The Y-peptides and Z-peptides (A1-Z and A2-Z) are derived from areas which, in human analogs, have
30 been shown to be recognized by T-cells, and may thus function in those analogs to elicit a T-helper cell response and/or a cytolytic T-cell response (Lamb, J.R., et al, Nature (1982) 300:66-69; Wabuke-Bunoti, M.A.N., et al, J Immunol (1984) 133:2194-2201). Both Z-peptides

and the N-peptide are derived from portions which correspond, in the human hemagglutinin, to areas of antigenic stability.

5 The foregoing peptides were used to immunize rabbits or mice both as prepared and as modified to enhance presentation size in the injected subject. For size enhancement, peptides were either self-polymerized with glutaraldehyde or covalently linked to keyhole limpet hemocyanin (KLH) using
10 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide.

For polymerization with glutaraldehyde, 4-10 mg/ml peptide was incubated at 21°C for 2 hr in 50 mM NaHPO₄ (pH 7.5), 150 mM glutaraldehyde. Insoluble products resulted after one or more hours, and the
15 precipitate was resuspended and mixed with Freund's adjuvant for immunization.

For conjugation to KLH, peptide at 4-10 mg/ml was incubated at 4°C for 20 min in the presence of 1 mg/ml carbodiimide in distilled water, adjusted to pH
20 3.5 with 1 N HCl. KLH was added at 1 mg/ml and the pH of the solution was adjusted to 9.0 with 5 volumes of distilled water containing 1 mM NaOH. The mixture was incubated for 2 hr at 21°C and dialyzed against phosphate-buffered saline at 4°C for 16 hr. After
25 dialysis, the conjugated peptides were mixed with Freund's adjuvant for immunization.

D.3. Immunization with Synthetic Peptides Antigenicity

30 The synthetic peptides were shown to be antigenic in rabbits and mice by ELISA assay of sera obtained from animals injected with each synthetic peptide. The same synthetic peptide was used as the antigen in the ELISA assay of 1C.6.

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For rabbits, each animal received a total of two inoculations. The first inoculation was administered intranodally with complete Freund's adjuvant. The following booster at three weeks was
5 injected intramuscularly with incomplete Freund's adjuvant. The animals received 125 μ g of free peptide or polymerized peptide per dose, or 165 μ g of KLH peptide conjugate (20 peptide:1 KLH). The animals were bled from the ear vein before the first inoculation to
10 obtain preimmune serum, and ten days after each subsequent inoculation.

For mice, eight-week-old female Balb/C mice received two inoculations of the same preparations as were used for rabbits, except that the peptide content
15 in each dose was 20 μ g. The first inoculation was given intramuscularly with complete Freund's adjuvant; the second, three weeks later, in incomplete adjuvant. Animals were bled from the eye seven or eight days after each inoculation.

20 Table 1 shows the results for the A1-X and A1-Y peptides and their derivatives.

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0.001
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Table 1

		<u>Peptide</u>	<u>Method of Peptide Presentation</u>	<u>Anti-Peptide Antibody Titer</u>
5	Rabbit 2A1	Al-X	Free	*
	Rabbit 2A2	"	Free	1/20,480
	Rabbit 2B1	"	Polymerized	1/80
	Rabbit 2B2	"	Polymerized	1/80
	Rabbit 2C1	"	Conjugated to KLH	1/5120
	Rabbit 2C2	"	Conjugated to KLH	1/5120
	* Animal died.			
10	Mouse 2fA	Al-X	Free	1/1280
	Mouse 2fB	"	Free	1/1280
	Mouse 2pA	"	Polymerized	1/320
	Mouse 2pB	"	Polymerized	<1/20
	Mouse 2cA	"	Conjugated to KLH	1/81,920
	Mouse 2cB	"	Conjugated to KLH	1/81,920
15	Mouse 3fA	Al-Y	Free	1/320
	Mouse 3fB	"	Free	<1/20
	Mouse 3pA	"	Polymerized	1/80
	Mouse 3pB	"	Polymerized	1/80
	Mouse 3cA	"	Conjugated to KLH	1/320
	Mouse 3cB	"	Conjugated to KLH	1/320

20 These data indicate that the synthetic peptides of this invention will elicit moderate to high antibody titers when administered in free and/or size-enhanced form.

25 Use of the Synthetic Peptides as Immunogenic Boosters

 Eight-week-old Balb/C mice received single intramuscular injections of 100 μ l of Equicine (Bayvet, Miles Laboratories). Equicine is a killed virus vaccine containing both A1 and A2 EIV serotypes.

30 Three weeks later, the mice received either a second dose of Equicine or a mixture containing 15 μ g each of the 7 peptides prepared in WD.2 in incomplete Freund's adjuvant. Six days after the second inoculation, the animals were bled from the eye, and the sera were

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assayed for neutralizing activity against EIV-A1 virus in a plaque reduction assay.

The results are given in Table 4.

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Table 4

	<u>Animal</u>	<u>Boost</u>	<u>Plaque reduction titer (EIV-A1)</u>
	10A	Equicine	1/160
	10B	Equicine	1/80-1/160
10	11A	EIV peps	1/40
	11B	"	>1/320
	11C	"	>>1/320
	11D	"	1/20
	11E	"	>1/320
	11F	"	1/80

15 The results show that the EIV peptides were as, or more, effective in 4 of 6 animals than additional killed virus in maintaining the level of neutralizing plaque-reduction antibodies. The same results may be achieved when the titer against EIV-A2 virus is analyzed.

20 A second similar experiment was performed to determine whether EIV synthetic peptides could be used in an adjuvant other than incomplete Freund's to boost neutralizing titer after a single inoculation of Equicine. The adjuvant preparation selected is a form of Halvogen™, an adjuvant used in Equicine vaccine and
25 approved for equine usage. It has the composition 7% Tween 20, 3% Arlacel 20, 50% soybean oil, and 1% Carbopol in H₂O.

30 Five groups of three mice each were inoculated with a 100 µl dose of Equicine. Group I received two boosts of Equicine at three week intervals; Group II received two boosts of 15 µg each peptide (105 µg total) in incomplete Freund's adjuvant (IFA); Group III received two boosts of 15 µg each peptide in the above

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Halvogen™ adjuvant; Group IV received two boosts of 15 µg each peptide in aqueous solution (no adjuvant); Group V received no boosts. The mice were bled eight days after the second boost; the sera in each group were
5 pooled, heat treated, and assayed in the plaque reduction assay. The results are presented in Table 5.

Table 5

10	<u>Group</u>	<u>Boost</u>	<u>Plaque reduction titer (EIV-A1)</u>
	I	Equicine	1/640
	II	EIV peps in IFA	1/1280
	III	EIV peps in Halvogen	1/640
	IV	EIV peps in aqueous (no adjuvant)	1/320
15	V	None	1/160

The results show that EIV peps in IFA boosted plaque reduction titer after a single primary vaccine inoculation better than subsequent vaccine boosts. In addition, EIV peps in Halvogen boosted as well as
20 vaccine boosts although EIV peps without adjuvant did not. Pooled sera from groups I, II, and III all showed at least a fourfold rise in titer over pooled sera from animals receiving no boost (group V). A fourfold titer increase is generally regarded as "seroconversion" in
25 the evaluation of vaccine preparations. Similar results may be obtained when the plaque reduction titer against EIV-A2 is analyzed.

In a converse experiment, 8-week-old female Balb/C mice received a single intramuscular injection of
30 a mixture containing 15 µg of each of the above peptides in complete Freund's adjuvant, while controls received similar inoculations of a heterologous peptide derived from unrelated viral sources (bovine viral diarrhea virus, or BVDV). Three weeks later, the mice

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received 10^5 pfu of UV-irradiated EIV-A1 or EIV-A2 intramuscularly in incomplete Freund's adjuvant. The animals were bled six days after the second inoculation and the sera analyzed as above. The results are shown in Table 6.

Table 6

	<u>Animal no. and initial treatment</u>	<u>Plaque reduction titer (EIV-A1)</u>
10	7D (EIV peps)	1/40
	7A (EIV peps)	>1/10*
	8B (BVD peps)	<1/10
	8A (BVD peps)	<1/10
	9B -0-	<1/10

*Assay not carried beyond 1/10 dilution.

These results show that the EIV peptides can also specifically prime the immune system of an animal for a subsequent inoculation of whole virus whereas an injection of heterologous peptide does not. Similar results may be obtained against EIV-A2.

D.4. Preparation of Recombinant Vaccinia Viruses Bearing Equine Influenza Antigens

Methods described by Mackett, et al (supra), were used to insert H7, H3, N7, and N8 into vaccinia carrier.

Each cDNA was first subcloned into pGS20 which contains the 7.5 k vaccinia gene promoter with unique BamHI and SmaI sites immediately downstream for the insertion of the desired heterologous gene, and flanked by vaccinia thymidine kinase (TK) gene. The tk- gene permits homologous recombination into the vaccinia genome and affords a method of selection (generation of tk-viruses).

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The constructions of the intermediate pGS20 derivative vectors containing EIV inserts are shown in Figure 6 and are described below. The nucleotide numbering referred to in the text corresponds to that in Figures 1-4. In order to place the initiation codons of each EIV cDNA as close to the vaccinia promoter as possible and to prevent possible interference with processing, the cDNAs were prepared for insertion by removal of homopolymeric dC or dG tracts that were generated during cDNA cloning and by providing convenient restriction sites proximal to the 5' and 3' ends.

For H7, the H7 cDNA was cloned into the PstI site of M13mp7, thus generating a SalI site at the 3' end of the DNA. A BamHI site was placed 3 nucleotides before the ATG start codon by converting the nucleotide at position 40 from A to C using site-specific mutagenesis. The H7 coding sequence was removed as a BamHI/SalI (blunted) fragment and cloned into the BamHI/SmaI digested pGS20.

For H3, nucleotides 1-94 and 1720-1797 were removed from H3 cDNA by digestion with AccI at the 5' end (GTCTAC) and BglI at the 3' end (GCCAAAAGGC). The deleted sequences were replaced using synthetic oligomers which contained SmaI restriction sites 8 nucleotides 5' of the ATG and 3 nucleotides 3' to the H3 termination codon while retaining the same coding sequence as represented in Figure 2. The resulting cDNA was removed as a SmaI/SmaI fragment and cloned into the SmaI site of pGS20.

For N7, the full-length cDNA was cloned into pUC9 so that a unique SmaI site was positioned at the 3' end. The upstream portion of the sequence was excised by treating with HindIII and EcoRI and inserted into

3 Cys

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HindIII/EcoRI digested M13mp9. This upstream sequence was modified to create an NdeI site by conversion of nucleotides 56 from G to C and 58 from A to T using site-specific mutagenesis. The heteroduplex phage was
5 digested with HindIII and EcoRI and the heteroduplex fragment was amplified in pBR322, then removed by treating with NdeI, blunting with Klenow, and digesting with EcoRI. The mutated excised fragment was ligated into SmaI/EcoRI digested pUC19. The SmaI site from
10 pUC19 is destroyed by the ligation and a unique BamHI site 5' to the N7 ATG results. The modified N7 5' fragment was removed from pUC19 as a BamHI/EcoRI fragment and religated to the remainder of the N7 sequence contained in pUC9 at the unique EcoRI site.
15 The desired sequence containing the entire N7 coding region was obtained as a BamHI/SmaI fragment and ligated into BamI/SmaI-digested pGS20.

For N8, the cDNA was cloned into the PstI site of pUC8, which placed a unique SmaI site 3' of the
20 cDNA. A unique AhaIII site already exists in N8 3 nucleotides 5' of ATG, and hence an AhaIII/SmaI digest generates a full-length cDNA which is blunt-end ligated into the SmaI site of pGS20.

The four vaccinia-EIV vectors described above
25 are designated pGS20-H7, pGS20-H3, pGS20-N7, and pGS20-N8, or, collectively, pGS20-EIV. Figure 6 shows the nucleotide sequence surrounding the coding region of each EIV cDNA inserted into the BamHI/SmaI cloning site of pGS20. The dotted line after each ATG represents
30 the sequence described in Figures 1-4 to the terminal PstI cloning sites (after the poly-G tracts) with the exception of H3, which was modified as described in the text.


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These recombinant vectors are transfected into cells infected with wild-type vaccinia virus, which was purchased from Wyeth Laboratories, Inc. (Marietta, PA) and plaque purified twice in CV-1 cells. A small
5 aliquot of virus stock (0.1 ml) is diluted with an equal volume of trypsin (0.25 mg/ml) and incubated for 30 min at 37°C with vortexing, followed by sonication to disperse any cell clumps. The virus is diluted to a concentration of 5×10^4 plaque-forming units per ml in
10 phosphate-buffered saline (PBS) containing 0.2% bovine serum albumin (BSA) and penicillin/streptomycin. CV-1 cells are infected in monolayers on 60 mm plates with 1 ml vaccinia virus to give a multiplicity of infection of 0.05 pfu/cell. The virus inoculum is incubated on the
15 cells for 2 hr at 37°C with rocking.

The DNA for transformation is prepared as described in Graham, et al. Virology (1973) 52:456; Stow, et al. J Gen Virol (1976) 33:447; and Frost, et al. Virology (1978) 91:39. Briefly, 5-10 µg of
20 plasmid (pGS20 derivative) DNA, and 1-2 µg wild-type vaccinia virus DNA are added to 1 ml of Hepes-buffered saline (0.14 molar NaCl, 5 mM KCl, 1 mM Na phosphate, 0.1% dextrose, 20 mM Hepes, pH 7.05), and 50 µl of 2.5 molar CaCl_2 is added. The solutions are mixed and
25 left at room temperature for 30 min, and the desired precipitate of DNA forms in this time.

The virus inoculum is aspirated from the CV-1 cell layers, and 1 ml of the DNA precipitate is substituted and the layers left at room temperature for
30 min, after which 9 ml prewarmed Eagle's MEM containing 8% FBS is added. The layers are incubated for 3.5 hr at 37°C, before aspirating off the medium and replacing it with 10 ml fresh Eagle's MEM containing 8% FBS. The monolayers are then left at 37°C for 2 days.



resulting in the development of vaccinia cytopathic effects. The cells and virus are harvested by scraping, spun down, and resuspended in 0.5 ml Eagle's MEM and frozen at -20°C.

5 The resuspended virus/cell pellets are freeze-thawed three times by freezing the cells at -20°C for 30 minutes and quickly thawing them at 37°C, followed by sonication for one minute to disperse the virus/cell clumps. The resulting crude virus stock is
10 then inoculated in serial 10-fold dilutions onto 143 cells (Mackett, M., et al, J Virol (1984) 49:857-864), a human tk- cell line. After two hours at 37°C, the 143 monolayers are then overlaid with 1% agarose containing 1x modified Eagle's medium, 5% fetal bovine serum, and
15 25 µg/ml 5-bromodeoxyuridine (BUdR). After incubation for two days at 37°C, BUdR-resistant plaques are picked and grown for 48 hours at 37°C in 24 well plates of 143 cell monolayers in the presence of 25 µg/ml BUdR.

 Virus is harvested and assayed for the presence
20 of the EIV-derived gene by DNA-DNA dot blot hybridization, as follows: cells are scraped from the dish into an Eppendorf centrifuge tube, centrifuged for 1 minute, and the cell pellet resuspended in 0.2 ml PBS. After freeze-thawing 3 times and sonicating as
25 described above, the sonicate is applied to a nitrocellulose filter and air dried. A wild-type virus control is also spotted onto the same filter. The filter is then placed on paper soaked in (1) 0.5 M NaOH, (2) 1 M Tris-HCl, pH 7.5, and (3) 2x SSC, for 5 minutes
30 each. The filter is then baked at 80°C under vacuum for 2 hours. The baked filter is prehybridized for 1-4 hours at 42°C in 5 ml 50% formamide, 4x SSC, 5x Denhardt's solution, and 0.1 mg/ml sheared and boiled salmon sperm DNA. The filter is then hybridized

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overnight at 42°C to 2×10^7 cpm of ^{32}P -labeled EIV gene probe in 5 ml prehybridization buffer. The probe is prepared by isolating 5 µg of EIV gene insert from one of the pGS20-derivative vectors and nick translating the DNA using a commercially available nick translation kit. The filter is then washed twice for 30 minutes in 0.5x SSC, 0.1% SDS, air dried, and autoradiographed overnight at -70°C.

Virus containing EIV gene inserts by dot blot hybridization are infected onto monolayers of 143 cells and left until a confluent cytopathic effect is obtained. The culture medium is then aspirated off and the cells lysed in 1% SDS, 0.1 M β-mercaptoethanol, 50 mM Tris-HCl, pH 7.8. The lysate is made 0.5 mg/ml in proteinase K, incubated 4 hours at 37°C, phenol extracted, ethanol precipitated, and analyzed by restriction enzyme analysis to show that the vaccinia genomes indeed contain the desired EIV genes.

The recombinant vaccinia so generated were designated vac-H7, vac-H3, vac-N7 and vac-N8 or collectively, vac-EIV.

D.5 Expression of EIV Surface Antigens by Vaccinia

Recombinants

Each vac-EIV recombinant was plaqued separately onto CV1 monolayers. The monolayers were washed, fixed, and incubated with rabbit anti-EIV-A1 (for vac-H7 and vac-N7) and rabbit anti-EIV-A2 (for vac-H3 and vac-N8). Antibody binding specifically to EIV surface antigens was detected by a subsequent incubation with ^{125}I -labeled staphylococcal A protein followed by autoradiography. In each case, all recombinant virus plaques bound the appropriate antibody.

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Immunoprecipitation of ^{35}S -labeled cellular lysates obtained from monolayers of CV1 cells infected by individual vac-EIV recombinants showed the presence of EIV protein. In the case of each recombinant virus, antisera raised against the appropriate whole virus immunoprecipitated a characteristic hemagglutinin or neuraminidase molecule identified by polyacrylamide gel electrophoresis and autoradiography.

10 D.6 Amplification of Vaccinia Recombinants

After the vac-EIV recombinants were identified as both carrying the appropriate EIV gene and expressing the encoded EIV protein, a single plaque of each is isolated x2 in 143 cells in the presence of BUdR. The twice purified isolate is amplified once in 143 cells in the presence of BUdR and subsequently in CV-1 cells in the absence of selective pressure. The resulting vac-EIV recombinants are stable genetically and can be further amplified in CV-1 cells.

20

D.7 Bioassay of Vaccinia Recombinants

Horses are used as subjects to assess the ability of the recombinant vaccinia to raise titers of neutralizing anti-EIV antibodies in serum.

25 In general, pairs of horses are inoculated with wild-type or recombinant virus by intradermal administration of $1-2 \times 10^8$ plaque-forming units distributed in 2-3 sites on the back, and are bled at days 0, 14, and 28. The sera are then tested for the presence of anti-EIV neutralizing antibodies in the plaque reduction assay of WC.7.

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In one experiment, four horses, 1-3 years old, housed in a P3 facility at Washington State University, Pullman, Washington, were ascertained to be serum

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negative with respect to anti-EIV antibodies, but were otherwise undocumented with respect to exposure.

Control and test virus were administered intradermally by scarification with a needle to bleeding at four
 5 locations on the animals' necks. Each animal received a total of 10^8 pfu of each indicated virus in a total of 0.4 ml (0.1 ml at each scar). VSV was used as a control; test viruses were H7, H3, N7, and N8, as described above (¶D.4).

10 The animals were bled at day 0 (preimmune), and at days 7, 14, and 21, and their sera assessed for neutralizing antibodies against each of EIV-A1 and EIV-A2 using the plaque reduction assay described in ¶C.7 hereinabove. The results, shown below in Table 7,
 15 indicate that while control VSV was not able to raise antiserum against EIV, all combinations of recombinant vaccinia containing DNA encoding H and/or N peptides, were able to do so.

Table 7

	<u>Horse</u>	<u>Vaccine</u>	<u>EIV Strain</u>	<u>Day</u>			
				0	7	14	21
25	Lola	VSV	A1	1/10	1/10	1/10	1/10
			A2	<1/10	<1/10	<1/10	<1/10
	Howard	N7,N8	A1	1/10	1/20	1/160	1/160
			A2	<1/10	1/40	1/160	1/160
	Fanny	H7,H3	A1	<1/10	1/160	1/320	1/160
		N7,N8	A2	<1/10	1/320	1/640	1/320
30	Leo	H7,H3	A1	<1/10	1/800	1/1600	1/800
			A2	<1/10	1/640	1/1280	1/640

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The results are maximum serum dilutions able to reduce EIV pfu by 50%. The results for Leo at days 7, 14, and 21 are based on an initial serum dilution of 1/50 rather than the 1/10 dilution used for all other results.

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Claims

1. A recombinant DNA sequence encoding a protein selected from the group consisting of equine hemagglutinin H7, equine hemagglutinin H3, equine neuraminidase N7, and equine neuraminidase N8.

2. A vaccine effective against influenza in equine hosts which comprises at least one DNA sequence of claim 1 or an immunologically effective portion thereof recombinantly integrated into vaccinia virus.

3. A method of immunizing horses against influenza which comprises administering the vaccine of claim 2.

4. A peptide useful in preparing a vaccine against equine influenza which comprises an amino acid sequence selected from the group consisting of:

Ala-Cys-Arg-Arg-Ser-Arg-Ser-Ser-Phe-Tyr-Ala-Glu-Met-Lys-Trp-Leu-Leu-Ser-Asn-Thr-Asp-Asn-Gly-Val-Phe-Pro-Gln-Met;

Cys-Lys-Arg-Glu-Pro-Ala-Leu-Ile-Ile-Trp-Gly-Ile-His-His-Ser-Gly-Ser-Thr-Ala-Glu-Gln-Thr-Arg-Leu-Tyr-Gly-Ser-Gly-Asn-Lys-Cys;

Arg-Ala-Ile-Gly-Lys-Cys-Pro-Arg-Tyr-Val-Lys-Gln-Lys-Ser-Leu-Met-Leu-Ala-Thr-Gly-Met-Lys-Asn-Val-Pro-Glu-Asn-Ser-Thr;

Thr-Gly-Val-Thr-Gln-Asn-Gly-Arg-Ser-Gly-Ala-Cys-Arg-Arg-Gly-Ser-Ala-Ser-Arg;

Lys-Leu-Tyr-Ile-Trp-Gly-Ile-His-His-Pro-Ser-Thr-Asn-Asn-Glu-Gln-Thr-Lys-Leu-Tyr-Ile-Gln-Glu-Ser-Gly-Arg-Val-Thr;

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Lys-Tyr-Ile-Lys-Gln-Asn-Thr-Leu-Lys-Leu-Ala-Thr-Gly-Met-Arg-Asn-Val-Pro-Glu-Lys-Gln-Ile; and
Gly-Ile-Phe-Gly-Ala-Ile-Ala-Gly-Phe-Ile-Glu-Asn-Gly-Lys-Lys.

5

5. A vaccine for immunizing horses against influenza wherein the peptide of claim 4 is optionally linked to a neutral carrier or polymerized.

10

6. A method of immunizing horses against influenza wherein the protocol is selected from boosting subject horses previously administered another vaccine against EIV with a vaccine comprising at least one peptide of claim 4; and

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administering a primer dose of a vaccine comprising at least one peptide of claim 4 to horses which are subsequently administered another vaccine against EIV..

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7. Recombinantly produced equine hemagglutinin H3, equine hemagglutinin H7, equine neuraminidase N7, or equine neuraminidase N8.

25

8. A vaccine effective for immunizing horses against influenza which comprises a protein selected from the group consisting of equine hemagglutinin H3, equine hemagglutinin H7, equine neuraminidase N7, and equine neuraminidase N8 or an immunologically effective portion thereof.

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9. A cDNA sequence substantially equivalent to a sequence selected from the group consisting of the equine H7 of Figure 1, the equine H3 of Figure 2, the equine N7 of Figure 3, and the equine N8 of Figure 4.

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10. A method for preparing a vaccine against equine influenza which comprises probing a cDNA library prepared from a fresh isolate of EIV from a desired animal with a probe derived from the cDNA sequence of claim 9.

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EIV - A1 hemagglutinin (M7)

ctgcagggggg gggggggggg ggggagcaaaa gcaaggggata caca

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ATG AAC ACT CAA ATT CTA ATA TTA GCC ATT TCG GCA TTC CTC TGT GTA CGT GCA GAT AAA ATC TGC CTA GGA CAT CAT GCT GTG TCT AAT
MET Aen Thr Gln Ile Leu Ile Leu Ala Ile Ser Ala Phe Leu Cys Val Arg Ala Asp Lys Ile Cys Leu Gly His His Ala Val Ser Aen 30

100
GCA ACC AAA CTA GAC ACC CTT ACT GAA AAG GGA ATA GAA GTT GTC AAT GCA ACA GAA ACA GTT GAA CAA AAA AAC ATC CCC AAG ATC TGC
Gly Thr Lys Val Asp Thr Leu Thr Glu Lys Gly Ile Glu Val Val Aen Ala Thr Glu Thr Val Glu Gln Lys Aen Ile Pro Lys Ile Cys 60

150
TCA AAA GGG AAA CAG ACT ATT GAC CTT GGT CAA TGT GGA TTA CTA GGG ACC ACT ATT GGT CCC CCC CAA TGC GAC CAA TTT CTT GAA TTC
Ser Lys Gly Lys Gln Thr Ile Asp Leu Gly Gln Cys Gly Leu Leu Gly Thr Thr Ile Gly Pro Pro Gln Cys Asp Gln Phe Leu Glu Phe 90

200
TCT GCT AAT TTA ATA ATT GAG AGA AGA GAA GGT GAT GAC ATT TGT TAT CCA GGC AAA TTT GAC AAT GAA GAA ACA TTC AGA CAA ATA CTC
Ser Ala Aen Leu Ile Ile Glu Arg Arg Glu Gly Asp Asp Ile Cys Tyr Pro Gly Lys Phe Asp Aen Glu Glu Thr Leu Arg Gln Ile Leu 120

250
AGA AAA TCC GGA CGA ATT AAA AAG GAG AAT ATG GGA TTC ACA TAT ACC GGA GTG AGA ACC GAT GGA GAG ACT AGC GCC TGT AGA AGG TCA
Arg Lys Ser Gly Gly Ile Lys Lys Glu Aen MET Gly Phe Thr Tyr Thr Gly Val Arg Thr Asp Gly Glu Thr Ser Ala Cys Arg Arg Ser 150

300
AGA TCT TCC TTT TAT GCA GAA ATG AAA TGG CTC CTA TCC AAC ACA GAC AAT GGG CTA TTC CCA CAA ATG ACA AAA TCC TAC AAG AAC ACT
Arg Ser Ser Phe Tyr Ala Glu MET Lys Trp Leu Leu Ser Aen Thr Asp Aen Gly Val Phe Pro Gln MET Thr Lys Ser Tyr Lys Aen Thr 180

350
AAG AGG GAG CCA GGT CTC ATA ATC TGG GGA ATC CAC CAC TCA GGA TCA ACC GCT GAA CAG ACT AGA TTG TAT GGA AGC CGA AAC AAG TTG
Lys Arg Glu Pro Ala Leu Ile Ile Trp Gly Ile His His Ser Gly Ser Thr Ala Glu Gln Thr Arg Leu Tyr Gly Ser Gly Aen Lys Leu 210

400
ATA ACA GTT TCG ACT TCC AAA TAC CAA CAA TCT TTT GCC CCA AGC CCT GGA CCA AGG CGG CAA ATA AAT GGC CAA TCA GGA AGA ATT GAC
Ile Thr Val Trp Ser Ser Lys Tyr Gln Gln Ser Phe Ala Pro Ser Pro Gly Pro Arg Pro Gln Ile Aen Gly Gln Ser Gly Arg Ile Aen 240

450
TTT TAC TGG CTG ATG TTA GAT CCC AAT GAT ACT GTT ACT TTT AAT CGC GCC TTT ATA GCA CCT GAC CGC GCC AGT TTT CTA AGA
Phe Tyr Trp Leu MET Leu Asp Pro Aen Asp Thr Val Thr Phe Ser Phe Aen Gly Ala Phe Ile Ala Pro Asp Arg Ala Ser Phe Leu Arg 270

500
GGT AAA TCT CTA GGA ATT CAG ACT GAC GCA CAA CTT GAC AAC AAT TGT GAA GGT GAA TGT TAT CAT ATT GGA GGT ACT ATA ATT AGC AAC
Gly Lys Ser Leu Gly Ile Gln Ser Asp Ala Gln Leu Asp Aen Aen Cys Glu Gly Glu Cys Tyr His Ile Gly Gly Thr Ile Ile Ser Aen 300

550
TTG CCC TTT CAA AAC ATT AAT AGC ACA GCA ATT GGG AAA TGC CCC AGA TAC GTA AAG CAA AAA AGC TTA ATG CTA GCA ACC GGA ATG AAA
Leu Pro Phe Gln Aen Ile Aen Ser Arg Ala Ile Gly Lys Cys Pro Arg Tyr Val Lys Gln Lys Ser Leu MET Leu Ala Thr Gly MET 330

600
AAT GTT CCT GAA AAT TCT ACA CAC AAA CAG TTA ACT CAT CAC ATG CGC AAA AAA AGA GGT TTA TTT GGT CCA ATA GCA GGA TTT ATT GAA
Aen Val Pro Glu Aen Ser Thr His Lys Gln Leu Thr His His MET Arg Lys Lys Arg Gly Leu Phe Gly Ala Ile Ala Gly Phe Ile Glu 360

650
AAT GGA TGG CAA GGA TTA ATA GAT GGA TGG TAT GGA TAC AGA CAT CAG AAT CCA CAA GGA GAA ACT GCT GCA GAC TAC AAA ACT ACA
Aen Gly Trp Glu Gly Leu Ile Asp Gly Trp Tyr Gly Tyr Arg His Gln Aen Ala Gln Gly Glu Gly Thr Ala Ala Asp Tyr Lys Ser Thr 390

700
CAA TCT GCT ATC AAT CAA ATA ACC GGG AAA TTA AAC AGA CTA ATA GAA AAA ACC AAC CAG CAA TTT GAA CTA ATA GAT AAT AAA TTC AAT
Gln Ser Ala Ile Aen Gln Ile Thr Gly Lys Leu Aen Arg Leu Ile Glu Lys Thr Aen Gln Gln Phe Glu Leu Ile Asp Aen Lys Phe Aen 420

750
CAA ATA GAA AAG CAA ATT GCC AAT GTT ATT AAC TGG ACT AGA GAT TCT ATC ATC CAA GTA TGG TCA TAT AAT GCA GAA TTC CTC GTG CCA
Glu Ile Glu Lys Gln Ile Gly Aen Val Ile Aen Trp Thr Arg Asp Ser Ile Ile Glu Val Trp Ser Tyr Aen Ala Glu Phe Leu Val Ala 450

800
GTG GAG AAT CAA CAC ACT ATT GAT TCA ACT GAT TCA GAG ATG AAC AAA TTA TAT GAA AAG GTA AGA AGA CAG CTG AGA CAA AAT GCT CAG
Val Glu Aen Gln His Thr Ile Asp Ser Thr Asp Ser Glu MET Aen Lys Leu Tyr Glu Lys Val Arg Arg Gln Leu Arg Glu Aen Ala Glu 480

850
GAA GAT GGT AAT GGC TGT TTT GAA ATA TTC CAC CAA TGT GAC AAT CAT TGC ATG GCC AGC ATT AGA AAC AAT ACA TAT CAT CAT AAA AAA
Glu Asp Gly Aen Gly Cys Phe Glu Ile Phe His Gln Cys Asp Aen Asp Cys MET Ala Ser Ile Arg Aen Aen Thr Tyr Asp His Lys Lys 510

900
TAC AGA AAG GAG CCA ATA CAA AAC AGA ATT CAG ATT GAT GCA CTA AAG TTG ACC ACC GGT TAC AAA GAT ATA ATA CTT TGG TTT AGC TTC
Tyr Arg Lys Glu Ala Ile Gln Aen Arg Ile Gln Ile Asp Ala Val Lys Leu Ser Ser Gly Tyr Lys Asp Ile Ile Leu Trp Phe Ser 540

950
GGG GCA TCA TGT TTC TTA TTT CTT GCC ATT GCA ATG GTT CTT GCT TTC ATA TGC ATA AAA AAT GGA AAC ATG CGG TGC ACT ATT TGT ATA TAA
Gly Ala Ser Cys Phe Leu Phe Leu Ala Ile Ala MET Val Leu Ala Phe Ile Cys Ile Lys Aen Gly Aen MET Arg Cys Thr Ile Cys Ile 570

1000
ggt tgaacaaaca ccccttcttt cctcttcccc cccccccccc ccccttcay

FIG. 1

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EIV - Al neuraminidase (N7)

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ctgcaggggg gggggggggg ggtttttttt tttttagcaa aagcagggta attttgaa

100

ATG AAT CCT AAT CAA AAA CTC TTT GCA TCA TCC GGA ATA GCA ATA GTG CTA GGA ATA ATA AAT CTT CTC ATA GGA ATA TCC AAT ATG AGT
MET Asn Pro Asn Gln Lys Leu Phe Ala Ser Ser Gly Ile Ala Ile Val Leu Gly Ile Ile Asn Leu Leu Ile Gly Ile Ser Asn MET Ser 10

200

TTA AAT ATA TCT CTA TAT TCA AAA GGG GAA AGC CAC AAG AAT AAT AAC CTA ACA TGC ACA AAT ATC AAC CAG AAT GAT ACC ACC ATG GTA
Leu Asn Ile Ser Leu Tyr Ser Lys Gly Glu Ser His Lys Asn Asn Asn Leu Thr Cys Thr Asn Ile Asn Gln Asn Asp Thr Thr MET Val 60

300

AAC AGC TAC ATC AAT AAC GCA ACA ATA ATT GAC AAA AGT ACA AAA ATA GAA AAC CCT GGT TAT CTA CTG CTG AAC AAA AGT CTA TGC AAC
Asn Thr Tyr Ile Asn Asn Ala Thr Ile Ile Asp Lys Ser Thr Lys Ile Glu Asn Pro Gly Tyr Leu Leu Leu Asn Lys Ser Leu Cys Asn 90

400

GTT GAA GGA TGG GTT GTA ATA GCA AAG GAC AAT GCG ATT AGA TTT GGA GAA AGC GAA CAA ATC ATA GTA ACT AGA GAA CCT TAT GTC TCA
Val Glu Gly Trp Val Val Ile Ala Lys Asp Asn Ala Ile Arg Phe Gly Glu Ser Glu Gln Ile Ile Val Thr Arg Glu Pro Tyr Val Ser 120

500

TGT GAT CCT CTA AGT TGC AAA ATG TAT GCT CTA CAC CAA GGT ACT ACA ATC AGA AAC AAG CAT TCA AAT AGT ACC ACA CAC GAC AGA ACA
Cys Asp Pro Leu Ser Cys Lys MET Tyr Ala Leu His Gln Gly Thr Thr Ile Arg Asn Lys His Ser Asn Ser Thr Thr His Asp Arg Thr 150

600

GCC TTC CGA GGG CTC ATT TCT ACT CCA TTA GGT AGC CCC CCA ACA GTG AGC AAC AGT GAA TTC ATA TGT GTT GGG TGG TCA AGC ACA AGC
Ala Phe Arg Gly Leu Ile Ser Thr Pro Leu Gly Ser Pro Pro Thr Val Ser Asn Ser Glu Phe Ile Cys Val Gly Trp Ser Ser Thr Ser 180

700

TGC CAT GAT GGG GTA AAC AGG ATG ACA ATT TGT GTA CAA GGA GAC AAT GAA AAT GCT ACT GCA ACA GTG TAT TAC AAC AAG AGA CTT ACA
Cys His Asp Gly Val Asn Arg MET Thr Ile Cys Val Gln Gly Asp Asn Glu Asn Ala Thr Ala Thr Val Tyr Tyr Asn Lys Arg Leu Thr 210

800

ACC ACT ATT AAA ACA TGG GCT AAA AAC ATT TTA AGA ACC CAA GAG TCT GAA TGT GTT TGT CAT AAC AGC ACT TGT GTA GTG GTA ATG ACT
Thr Thr Ile Lys Thr Trp Ala Lys Asn Ile Leu Arg Thr Gln Glu Ser Glu Cys Val Cys His Asn Ser Thr Cys Val Val Val MET Thr 240

900

GAT GGG CCC GCA AAT AAC CAG GCG TTC ACA AAA GTA ATA TAC TTT CAT AAA GGA ATG ATA ATA AAA GAA GAA TCA CTA AAA GGT TCA GCC
Asp Gly Pro Ala Asn Asn Gln Ala Phe Thr Lys Val Ile Tyr Phe His Lys Gly MET Ile Ile Lys Glu Glu Ser Leu Lys Gly Ser Ala 270

1000

AAA CAC ATA GAA GAA TGT TCT TGT TAT GGT CAT AAT CAA AGA GTG ACT TGT GTC TGC AGA GAC AAC TGG CAG GGT GCA AAT AGA CCT ATT
Lys His Ile Glu Glu Cys Ser Cys Tyr Gly His Asn Gln Arg Val Thr Cys Val Cys Arg Asp Asn Trp Gln Gly Ala Asn Arg Pro Ile 300

1100

ATA GAG ATT GAC ATG AAT AAA TTG GAA CAT ACA AGT AGA TAT ATA TGC ACA GCG GTA TTA ACA GAC ACC AGT AGA CCC AAG GAT AAA ACA
Ile Glu Ile Asp MET Asn Lys Leu Glu His Thr Ser Arg Tyr Ile Cys Thr Gly Val Leu Thr Asp Thr Ser Arg Pro Lys Asp Lys Thr 330

1200

ATA GGG GAA TGC TTC AAT CCT ATT ACT GGA AGC CCT GGT GCA CCA GGG ATA AAA GGT TTC GGA TTC CTA AAT GAG GAT AAT ACT TGG CTA
Ile Gly Glu Cys Phe Asn Pro Ile Thr Gly Ser Pro Gly Ala Pro Gly Ile Lys Gly Phe Gly Phe Leu Asn Glu Asp Asn Thr Trp Thr 360

1300

GGG AGA ACA ATC AGC CCC AGA TTG AGG AGT GGA TTT GAA ATG CTG AAG ATA CCT AAT GCT GGG ACT GAC CCA GAG TCC AAA ATA AAA GAA
Gly Arg Thr Ile Ser Pro Arg Leu Arg Ser Gly Phe Glu MET Leu Lys Ile Pro Asn Ala Gly Thr Asp Pro Glu Ser Lys Ile Lys Glu 390

1400

ACA CAA GAA ATA GTT AGT AAT GAC AAT TGG TCA GGC TAT TCC GGA AGT TTC ATT GAT TAT TGG AAT GAC AAC AGT GAA TGC TAC AAT CCA
Arg Gln Glu Ile Val Ser Asn Asp Asn Trp Ser Gly Tyr Ser Gly Ser Phe Ile Asp Tyr Trp Asn Asp Asn Ser Glu Cys Tyr Asn Pro 420

1500

TGT TTC TAT GTA GAA TTA ATT AGA GGA AGG CCT GAA GAA GCA AAA TAT GTT GAA TGG ACC AGT AAC AGC CTA ATT GCA CTA TGT GGG AGC
Cys Phe Tyr Val Glu Leu Ile Arg Gly Arg Pro Glu Glu Ala Lys Tyr Val Glu Trp Thr Ser Asn Ser Leu Ile Ala Leu Cys Gly Ser 450

1600

CCA ATC TCA GTT GGG TCT GGA TCT TTC CCT GAT GGG GCA CAA ATT AAA TAC TTT TGG TAA aacgaa aaaaaacccct gttttctactc cccccccccc
Pro Ile Ser Val Gly Ser Gly Ser Phe Pro Asp Gly Ala Gln Ile Lys Tyr Phe Ser 469

1700

ccccctgcaq

FIG. 3

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ZIV - A2 neuraminidase (N8)

ctgcagggggg gggggagcag gaqttttaa

50
 ATG AAT CCA AAT CAA AAG ATA ATA GCA ATT GGA TCT GCA TCA TTA GGA ATA CTA ATC CTC AAC GTC ATT CTC CAT GTA GTC AGC ATT ATA
 MET Aen Pro Aen Gln Lys Ile Ile Ala Ile Gly Ser Ala Ser Leu Gly Ile Leu Ile Leu Aen Val Ile Leu His Val Val Ser Ile Ile 30

100
 150
 GTA ACA GTA CTG GTC CTC AAT AAC AAT GGA ACA GGT CTG AAC TGC AAC GGG AGC ATC ATA AGA GAG TAC AAT GAA ACA GTA AGA GTA GAA
 Val Thr Val Leu Val Leu Aen Aen Aen Gly Thr Gly Leu Aen Cys Aen Gly Thr Ile Ile Arg Glu Tyr Aen Glu Thr Val Arg Val Glu 60

200
 250
 AGA ATT ACT CAA TGG TAT AAT ACT AAT ACA ATC GAG TAT ATA CAG AGA CCT TCA AAT GAA TAC TAC ATG AAC AAC ACC GAA CCA CTG TGT
 Arg Ile Thr Gln Trp Tyr Aen Thr Aen Thr Ile Glu Tyr Ile Glu Arg Pro Ser Aen Glu Tyr Tyr MET Aen Aen Thr Glu Pro Leu Cys 90

300
 350
 GAG GCC CAG GGC TTT GCA CCA TTT TCC AAA GAT AAT GGA ATA CGA ATT GCG TCG AGA GGT CAT GTT TTT GTA ATA AGA GAA CCT TTT GTC
 Glu Ala Gln Gly Phe Ala Pro Phe Ser Lys Asp Aen Gly Ile Arg Ile Gly Ser Arg Gly His Val Phe Val Ile Arg Glu Pro Phe Val 120

400
 450
 TCA TGT TCG CCC TTA GAA TGT AGA ACC TTT TTC CTC ACA CAG GGC TCA TTA CTT AAT GAC AAA CAT TCT AAC GGC ACA GTG AAG CAC CGA
 Ser Cys Ser Pro Leu Glu Cys Arg Thr Phe Phe Leu Thr Gln Gly Ser Leu Leu Aen Asp Lys His Ser Aen Gly Thr Val Lys Asp Arg 150

500
 550
 AGT CCA TAT AGG ACT TTG ATG AGT GTC AAA GTA GGG CAA TCA CCT AAT GTG TAT CAA GCT AGG TTT GAA TCG GTC GCA TGG TCA GCA ACA
 Ser Pro Tyr Arg Thr Leu MET Ser Val Lys Val Gly Gln Ser Pro Aen Val Tyr Gln Ala Arg Phe Glu Ser Val Ala Trp Ser Ala Thr 180

600
 650
 GCA TGC CAC GAT GGG AAA AAG TGG ATG ACA GTT GGA GTC ACA GGG CCC GAT AAT CAA GCA GTT GCA GTA GTG AAC TAT GGA GGT GTT CGG
 Ala Cys His Asp Gly Lys Lys Trp MET Thr Val Gly Val Thr Gly Pro Asp Aen Gln Ala Val Ala Val Val Aen Tyr Gly Gly Val Pro 210

700
 750
 GTT GAT ATC ATT AAT TCA TGG GCA GGG GAT ATC CTA AGA ACC CAA GAA TCG TCA TGC ACC TGC ATT AAA GGA GAC TGT TAT TGG GTG ATG
 Val Asp Ile Ile Aen Ser Trp Ala Gly Asp Ile Leu Arg Thr Gln Glu Ser Ser Cys Thr Cys Ile Lys Gly Asp Cys Tyr Trp Val MET 240

800
 850
 ACT GAT GGA CGG GCA AAC AGG CAA GCT AAA TAT AGG ATA TTC AAA GCA AAA GAT GGA AGA ATA ATT GGG CAG ACT GAT ATA AGT TTC AAT
 Thr Asp Gly Pro Ala Aen Arg Gln Ala Lys Tyr Arg Ile Phe Lys Ala Lys Asp Gly Arg Ile Ile Gly Gln Thr Asp Ile Ser Phe Aen 270

900
 950
 GGG GGA CAC ATA GAG GAG TGT TCT TGT TAC CCC AAT GAA GGG AAG GTG GAG TGT GTA TGC AGG GAC AAC TGG ACT GGA ACA AAT AGA CCA
 Gly Gly His Ile Glu Glu Cys Ser Cys Tyr Pro Aen Glu Gly Lys Val Glu Cys Val Cys Arg Asp Aen Trp Thr Gly Thr Aen Arg 300

1000
 1050
 ATT CTG GTA ATA TCT CCT GAT CTA TCG TAC ACA GTC GGA TAT TTG TGT GCT GGC ATT CCC ACT GAC ACT CCT AGG GGA GAG GAT AGT CAA
 Ile Leu Val Ile Ser Pro Asp Leu Ser Tyr Thr Val Gly Tyr Leu Cys Ala Gly Ile Pro Thr Asp Thr Pro Arg Gly Glu Asp Ser 330

1100
 1150
 TTC ACA GGC TCA TGC ACA AGC CCT TTG GGA AAT AAA GGA TAC GGT GTA AAG GGC TTC GGG TTT CGA CAA GGA AAT GAC GTA TGG CCC GGA
 Phe Thr Gly Ser Cys Thr Ser Pro Leu Gly Aen Lys Gly Tyr Gly Val Lys Gly Phe Gly Phe Arg Gln Gly Aen Asp Val Trp Ala Gly 360

1200
 1250
 AGG ACA ATT AGT AGG ACT TCG AGA TCA GGA TTC CAA ATA ATA AAA ATC AGG AAT GGT TGG ACA CAG AAC AGT AAA GAC CAA ATC AGA AAG
 Arg Thr Ile Ser Arg Thr Ser Arg Ser Gly Phe Glu Ile Ile Lys Ile Arg Aen Gly Trp Thr Gln Aen Ser Lys Asp Gln Ile Arg 390

1300
 1350
 CAA GTG ATT ATT GAT AAC CTA AAT TGG TCA GGA TAT AGC GGT TCT TTC ACA TTG CCG GTT GAA CTA ACA AAA AAA GGA TGT TTA GTC CCC
 Gln Val Ile Ile Asp Aen Leu Aen Trp Ser Gly Tyr Ser Gly Ser Phe Thr Leu Pro Val Glu Leu Thr Lys Lys Gly Cys Leu Val Pro 420

1400
 1450
 TGT TTC TGG GTT GAA ATG ATC AGA GGT AAA CCT GAA GAC ACA ACA ATA TGG ACC TCT AGC AGC TCC ATT GTG ATG TGT GGA GTA GAC CAT
 Cys Phe Trp Val Glu MET Ile Arg Gly Lys Pro Glu Asp Thr Thr Ile Trp Thr Ser Ser Ser Ser Ile Val MET Cys Gly Val Asp His 450

1481
 cccccctgcag

FIG. 4

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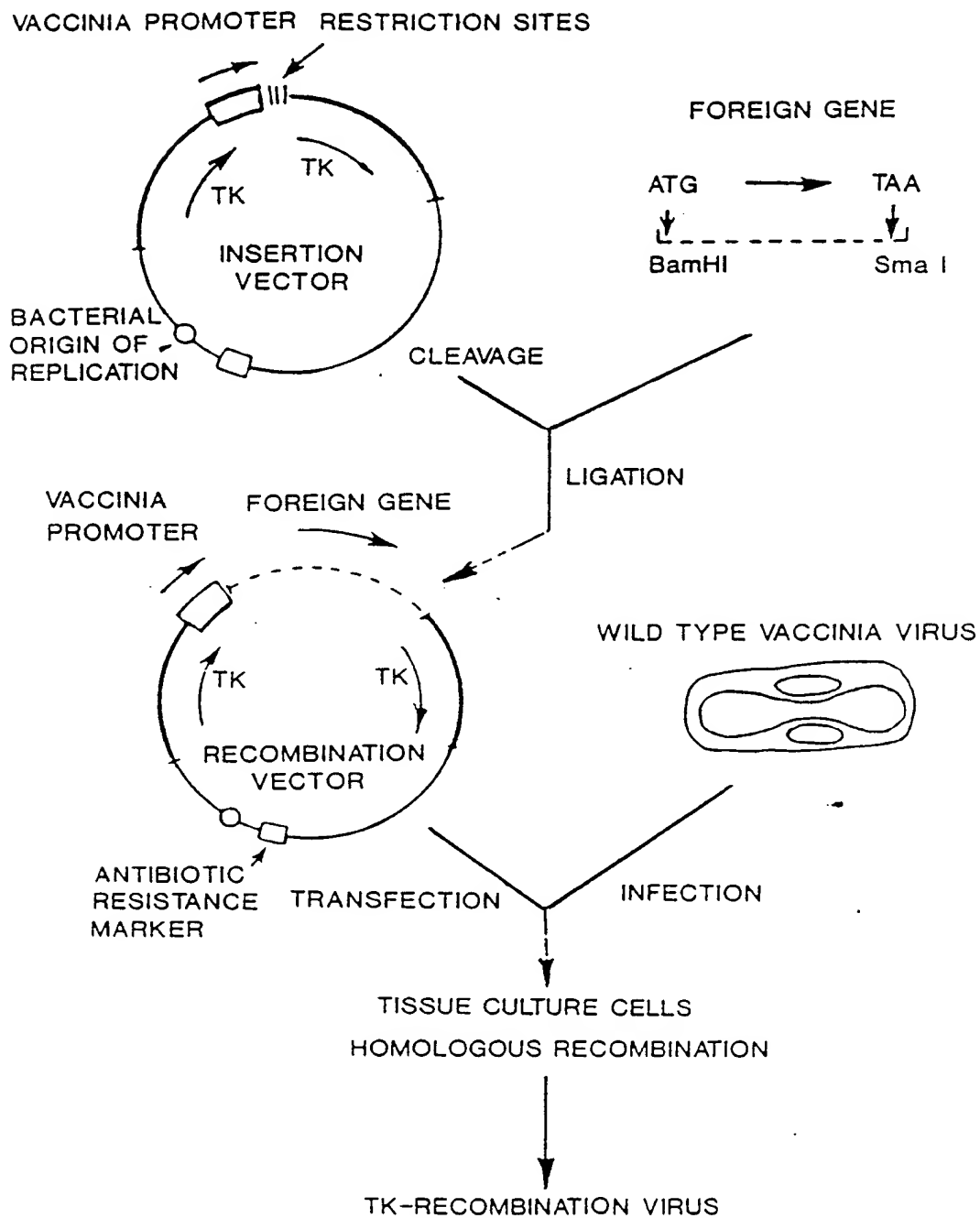


FIG. 5

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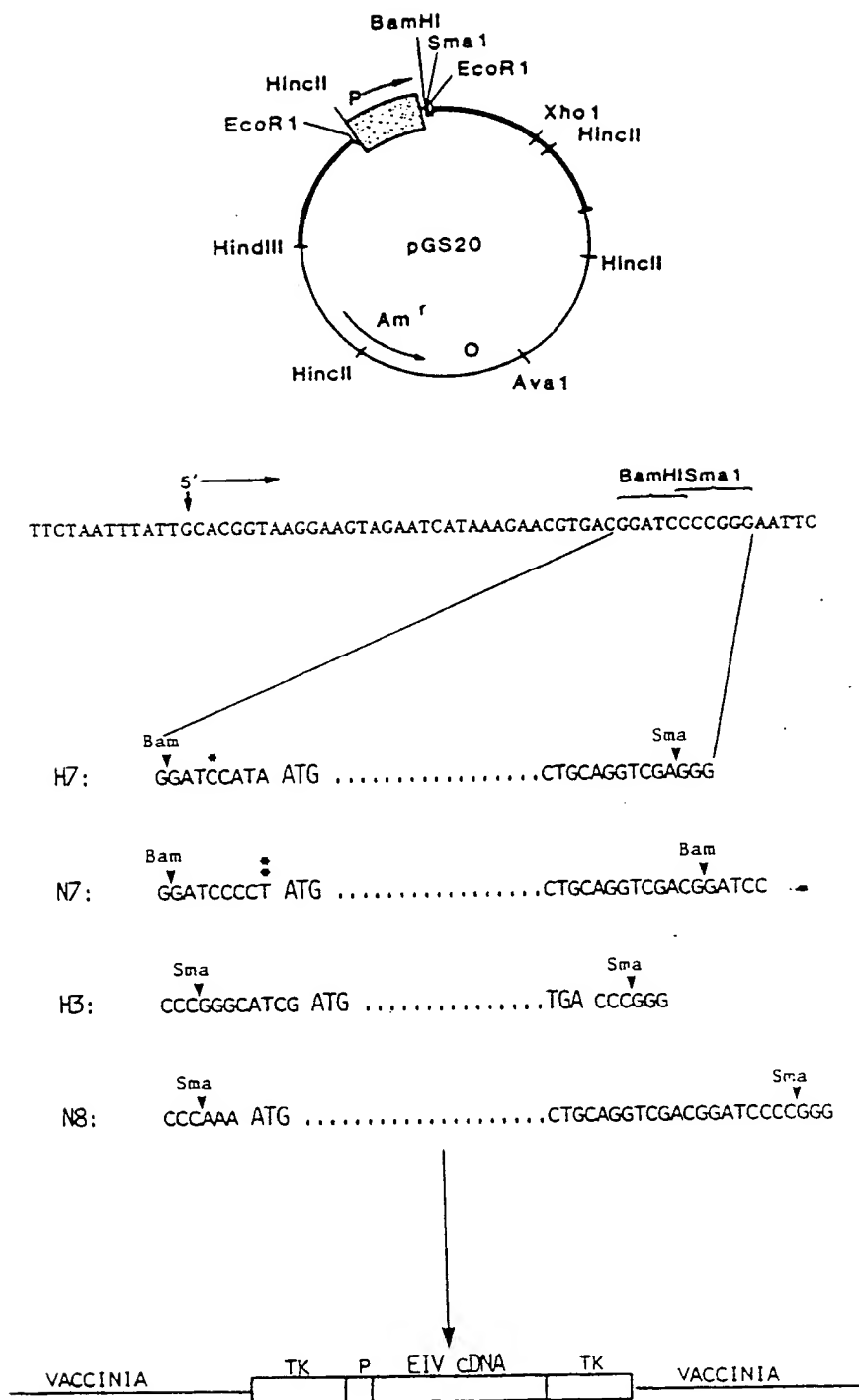


FIG. 6

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INTERNATIONAL SEARCH REPORT

International Application No PCT/US86/01343

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ³		
According to International Patent Classification (IPC) or to both National Classification and IPC		
U.S. CL. 536/27; 424/88; 530/324		
INT. CL. -4- CO7H 15/12; A61K 39/00; CO7K 7/10		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁴		
Classification System	Classification Symbols	
US	536/27; 424/88; 530/324	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁵		
III. DOCUMENTS CONSIDERED TO BE RELEVANT ¹⁴		
Category [*]	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
A	US, A, 4,357,421 Published 2, November 1982, (Emtage)	1-3, 5, 6, 8-10
A	US, A, 4,455,142 Published 19, June 1984, (Martins)	2, 3, 5, 6, 8
X, P	US, A, 4,537,769 Published 27, August 1985, (Cerini)	2, 3, 5, 6, 8
X, P	US, A, 4,548,814 Published 22, October 1985, (Rinehart, Jr.)	1-3, 5, 6, 8-10
A	US, A, 4,474,757 Published 2, October 1984, (Arnon)	2, 3, 5, 6, 8
A, P	US, A, 4,578,269 Published 25 March 1986, (Morein)	2, 3, 5, 6, 8
A	Ann. Rev. Microbiol. Published 1984, Vol. 3, pages 221-235, (Brown)	1-10
<div style="display: flex; justify-content: space-between;"> <div style="width: 48%;"> <p>[*] Special categories of cited documents: ¹⁵</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 48%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"Δ" document member of the same patent family</p> </div> </div>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search ²	Date of Mailing of this International Search Report ¹	
22 August 1986	02 SEP. 1986	
International Searching Authority ¹	Signature of Authorized Officer ¹⁰	
ISA/US	<i>Dellu H. Philler</i>	

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III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No ¹⁸
A	Journal of American Veterinarian Medical Association, Vol. 143, No. 6 pages 587-590, (Waddell), Published 15 Sept. 1963	1-10
A	Acta Virlogica, Vol. 2, pages 52-61, published 1958, (Sovinova)	1-10
A	Journal of Virology, Published March 1984, pages 857-864, Vol. 49, No. 3, (Mackett)	1-10
A	Proceeding National Academy of Science USA, Vol. 80, pages 7155-7159, Published Dec. 1983, (Smith)	1-10
A	Nature, Vol. 311, Published Sept. 1984, pages 65-69, (Bennink)	1-3,5,6, 8-10
A	Biotechniques, Published Nov./Dec. 1984, pages 306-312, (Smith)	2,3,5,6, 8

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FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. ☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE ¹⁰

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☐ Claim numbers because they relate to subject matter ¹² not required to be searched by this Authority, namely:

2. ☐ Claim numbers because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out ¹³, specifically:

VI. ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING ¹⁴

This International Searching Authority found multiple inventions in this international application as follows:

I. Claims 1, 9 and 10 drawn to DNA sequences classified in U.S. Class 536.

II. Claims 2, 3, 5, 6 and 8 drawn to compositions and methods of vaccination classified in U.S. Class 424 Subclass 88.

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.

2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

☐ The additional search fees were accompanied by applicant's protest.

☒ No protest accompanied the payment of additional search fees.

PCT/US86/01343

III. Claims 4 and 7 drawn to peptides classified in U.S.
Class 530 Subclass 324.

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